

GenCore version 5.1.6  
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OM nucleic - nucleic search, using sw model

Run on: October 25, 2005, 22:08:33 ; Search time 5981 Seconds  
(without alignments)

11341.001 Million cell updates/sec

Title: US-10-070-255-4

Perfect score: 1782

Sequence: 1 atgagcgatcacagacaa.....taaatgtcttcattccttg 1782

Scoring table:

IDENTITY NUC

Gapop 10.0, Gapext 1.0

Searched: 34239544 seqs, 19032134700 residues

Total number of hits satisfying chosen parameters: 69479088

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST:\*

1: gb\_est1:\*

2: gb\_est2:\*

3: gb\_hc:\*

4: gb\_est3:\*

5: gb\_est4:\*

6: gb\_est5:\*

7: gb\_est6:\*

8: gb\_gss1:\*

9: gb\_gss2:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	821	46.1	1981	3	BC024293 Homo sapi
2	653.8	36.7	998	5	BQ064251 AGENCOURT
3	651.6	36.6	990	5	BQ060776 AGENCOURT
4	650	36.5	691	4	BQ668241 UI-E-CKI-
5	648.2	36.4	970	5	BQ065026 AGENCOURT
6	639.8	35.9	730	1	AU122438
7	634.2	35.6	829	4	BG757448
8	627.8	35.2	835	4	BG758677
9	627.8	35.2	1022	5	BQ057616 AGENCOURT
10	624.6	35.1	843	4	BG758425
11	624.4	35.0	892	5	BQ057008
12	598	33.6	609	4	BM692924
13	574.8	32.3	813	4	BG684262
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15	534.6	30.0	626	5	BQ184171
16	521.6	29.3	620	5	BQ185857
17	513	28.8	769	4	BG684980
18	505.6	28.4	782	4	BG759163
19	500.6	28.1	914	4	BG340557
20	494.6	27.8	951	4	BG685372
21	491.8	27.6	1064	5	BQ065934
22	480.4	27.5	744	9	AY39018
23	478.8	26.9	737	4	BF975008
24	478.2	26.8	1075	5	BQ058038 AGENCOURT

25	471	26.4	1051	5	BQ061744
c	462.6	26.0	855	5	BX760749
26	461.2	25.9	575	2	AW974284
27	455.4	25.6	822	4	BM478075
28	453.2	25.4	965	4	BG757633
29	450.4	25.3	504	4	BM697878
c	439	24.6	439	1	AI136989
30	428.6	24.1	501	1	AL135642
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35	395	22.2	744	9	AY399020
36	384.8	21.6	430	2	AW403206
37	382.4	21.5	850	5	BQ642454
38	382	21.4	506	5	BX283339
39	373.4	21.0	522	2	AW044949
40	370.8	20.8	520	2	AW503340
41	355	19.9	478	5	BX283759
42	354.6	19.9	847	7	CR579211
43	342.4	19.2	646	1	AL965834
44	329.6	18.5	498	4	BG759800
45	326.4	18.3	658	2	BB623950

## ALIGNMENTS

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LOCUS BC024293 1981 bp mRNA linear HTC 04-MAR-2003  
DEFINITION Homo sapiens, clone IMAGE:5088007, mRNA.  
ACCESSION BC024293  
VERSION BC024293.1 GI:22028130  
KEYWORDS HTC.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
REFERENCE 1 (bases 1 to 1981)  
AUTHORS Strausberg,R.  
TITLE Direct Submission  
JOURNAL Submitted (26-FEB-2002) National Institutes of Health, Mammalian Gene Collection (MGC), Cancer Genomics Office, National Cancer Institute, 31 Center Drive, Room 11A03, Bethesda, MD 20892-2590, USA  
REMARK NIH-MGC Project URL: <http://mgc.nci.nih.gov>  
COMMENT Contact: MGC help desk  
Email: [cgaps-remail.nih.gov](mailto:cgaps-remail.nih.gov)  
Tissue Procurement: Lou Staudt  
cDNA Library Preparation: Rubin Laboratory  
DNA Library Arrayed by: The I.M.A.G.E. Consortium (ILLNL)  
BC Cancer Agency, Vancouver, BC, Canada  
info@bcgsc.bc.ca  
Steven Jones, Jennifer Asano, Ian Bosdet, Yaron Butterfield, Susanna Chan, Readman Chiu, Chris Fjell, Erin Garland, Ran Guin, Letticia Hsiao, Martin Krzywinski, Reta Kutsche, Oliver Lee, Soo Sen Lee, Victor Ling, Carrie Mathewson, Candice McLeavy, Steven Nees, Pawan Pandoh, Anna-Lisa Prabhu, Parvaneh Saeedi, Jacqueline Schein, Duane Smalus, Michael Smith, Lorraine Spence, Jeff Scott, Michael Thorne, Miranada Tsai, Natasja van den Bosch, Jill Vardy, George Yang, Scott Zuyderduyn, Marco Marra.

Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/ILLNL at: <http://image.llnl.gov>  
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This clone has the following problem: retained intron.

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/clone_lib="NIH_MGC_99"
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/notes="Vector: pOTB7"

DEFINITION
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BQ064251
VERSION
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KEYWORDS
EST.
SOURCE
Homo sapiens (human)
ORGANISM
Homo sapiens
REFERENCE
1 (bases 1 to 998)
NIH-MGC http://mgc.nci.nih.gov/.
AUTHORS
National Institutes of Health, Mammalian Gene Collection (MGC)
TITLE
Unpublished (1999)
JOURNAL
COMMENT
Contact: Robert Strausberg, Ph.D.
Email: c9apbs-r@mail.nih.gov
Tissue Procurement: Lou Staudt
cDNA Library Preparation: Rubin Laboratory
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
DNA Sequencing by: Agencourt Bioscience Corporation
Clone distribution: MGC clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
http://image.llnl.gov
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/notes="Organ: lymph; Vector: pOTB7; Site: 1: XhoI; Site 2:
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into EcoRI/XhoI sites using the following 5' adaptor:
GGCAGCAG(G). Size-selected >500bp for average insert size
1.8kb. Library constructed by Ling Hong in the laboratory
of Gerald M. Rubin (University of California, Berkeley)
using ZAP-cDNA synthesis kit (Stratagene) and Superscript
II RT (Life Technologies). Note: this is a NIH_MGC
Library."

ORIGIN
Query Match 36.7%; Score 653.8; DB 5; Length 998;
Best Local Similarity 97.0%; Pred. No. 5.9e-165;
Matches 617; Conservative 0; Mismatches 18; Indels 3; Gaps 1;

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QY 92 AGTACTTTTATGAAGACTGGTCTTTGTGATGGATGAAGAGAGTCCAGTATGCTTCCT 151
DB |||||||
QY 487 ACCATGGCAGCAGGTCTGAATCCATCTCTTTGCGATTAAACATCGACAAACAAGGATTG 546
DB |||||||
QY 152 ACCATGGCAGCAGGTCCGAATCCATCTCTTTGCGATTAAACATCGACAAACAAGGATTG 211
DB |||||||
QY 547 AACGGCAGAGTAGTTTGTCTCCACCGTTTCAGACCTCTTAAAGAGTCAACGCGAAC 606
DB |||||||
QY 212 AACGGCAGAGTAGTTTGTCTCCACCGTTTCAGACCTCTTAAAGAGTCAACGCGAAC 271
DB |||||||
QY 607 GTGACCTCTTGTGAAGGAGTCCACGCAAGGAGTGGAGCAGCCTGTTCAGGGAGATCACA 666
DB |||||||
QY 272 GTGA---CCTTGTGAAGGAGTCCACGCAAGGAGTGGAGCAGTGTTCAGGGAGATCACA 328
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DB |||||||
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DB |||||||
QY 727 GTCTCCAGGAATGTGAGTGTGATGCCAATGCAAAAGAGCGGAGAGAAAGAAACAAA 786
DB |||||||
QY 389 GTGTCCAGGAATGTGAGTGTGATGCCAATGCAAAAGAGCGGAGAGAAAGAAACAAA 448
DB |||||||
QY 787 GTGACCAACATAATCTCATTTTGTATGATGAGGAAAGATGAGCAGAACTCTTGGGACGTGTTT 846
DB |||||||


```



375 Newton Road, 4156 MEBRF, Iowa City, IA 52242, USA

Tel: 319 335 8250

Fax: 319 335 9565

Email: bento-soares@iowa.edu

Tissue Procurement: Dr. Gregg Hageman

cDNA Library preparation: Dr. M. Bento Soares, University of Iowa

cDNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa

DNA Sequencing by: Dr. M. Bento Soares, University of Iowa

Clone Distribution: Researchers may obtain clones from Research

Genetics (www.resgen.com).

Seq primer: M13 Forward

POLYA=Yes.

Location/Qualifiers

#### FEATURES

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/notes="Organ: eye; Vector: pT7T3-Pac (Pharmacia) with a modified polylinker; Site 1: EcoR I; Site 2: Not I; UI-E-CK1 is a normalized cDNA library containing the following tissue(s): Retina Foveal and Macular. The library was constructed according to Bonaldo, Lennon and Soares, Genome Research, 6:791-806, 1996. First strand cDNA synthesis was primed with an oligo-dT primer containing a Not I site. Double stranded cDNA was ligated to an EcoR I adaptor, digested with Not I, and cloned directionally into pT7T3-Pac vector. The oligonucleotide used to prime the synthesis of first-strand cDNA contains a library tag sequence that is located between the Not I site and the (dT)18 tail. The sequence tag for this library is GACC. This library was created for the program, Gene discovery in the Visual System, supported by National Eye Institute (NEI).  
TAG\_TISSUE=Foveal and Macular Retina  
TAG\_LIB=UI-E-CK1  
TAG\_SEQ=GTCC"

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QY 1170 TCCGCTGAAGGTGCTGCACAAATGATCCCGACATCCTCTTCCCTGTGTCAGTGGGTGGGCTC 1229  
DB 631 TCCGCTGAAGGTGCTGCACAAATGATCCCGACATCCTCTTCCCTGTGTCAGTGGGTGGGCTC 572  
QY 1230 CTACAGCCAGCAGATGCCCCCTCGGAAGCTGGGAAGCGGACAGGACGAGGACCA 1289  
DB 571 CTACAGCCAGCAGATGCCCCCTCGGAAGCTGGGAAGCGGACAGGACGAGGACCA 512  
QY 1290 CGTTCTCCCGGATCCTGGACTTCGGTACAGTGTGGAAGCCAGCTCTCCAGGCCACCGAAG 1349  
DB 511 CGTTCTCCCGGATCCTGGACTTCGGTACAGTGTGGAAGCCAGCTCTCCAGGCCACCGAAG 452  
QY 1350 TCCTCTGAGCAGCTGTTACCTTCTGCTCAGTGCAGGATCCATGACATAGTAGTGAAT 1409  
DB 451 TCCTCTGAGCAGCTGTTACCTTCTGCTCAGTGCAGGATCCATGACATAGTAGTGAAT 392  
QY 1410 GCCCAGGCCACGTGCGCATGATGAACAGGAGGATGAGCTGGAGGAGGACAGATC 1469  
DB 391 GCCCAGGCCACGTGCGCATGATGAACAGGAGGATGAGCTGGAGGAGGACAGATC 332  
QY 1470 ACTGCGAAACCTGCTCGACCGTGAGATGGAGCACTCAGCCGCGCTCCGGAAGAGGTGGA 1529  
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DB 211 GCCCAGCTATCTTCTGCTATTTTGTGAGGAGATCTTAACCCACGTCGAGAACCATGTGGTG 152  
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DB 92 CCAGTCTTTCTGAGAGCTGTGTTTCTCTGAGACTTTTCTATGTATGTATGTAGCCCAATAAATTT 33  
QY 1769 GCTTTTCATTCCTTG 1782  
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EST.  
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ORGANISM  
Homo sapiens  
REFERENCE  
1 (bases 1 to 970)  
NTH-MGC http://mgc.nci.nih.gov/  
National Institutes of Health, Mammalian Gene Collection (MGC)  
Unpublished (1999)  
Contact: Robert Strausberg, Ph.D.  
Email: cgapbs-romail.nih.gov  
Tissue Procurement: Lou Staudt  
cDNA Library Preparation: Rubin Laboratory  
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
DNA Sequencing by: Agencourt Bioscience Corporation  
Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:  
http://image.llnl.gov  
plate: LLCM2107 row: 1 column: 18  
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ORIGIN  
Query Match 36.4%; Score 648.2; DB 5; Length 970;  
Best Local Similarity 97.0%; Pred. No. 1.9e-163;  
Matches 672; Conservative 0; Mismatches 18; Indels 3; Gaps 1;

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LOCUS  
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VERSION AUI22438.1 GI:10937708  
KEYWORDS EST  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Eukaryote; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
REFERENCE 1 (bases 1 to 730)  
AUTHORS Ota, T., Nishikawa, T., Suzuki, Y., Ishii, S., Saito, K., Kawai, Y.,  
Yanamoto, J., Wakamatsu, A., Nakamura, Y., Nagai, T., Sugano, S. and  
Isogai, T.  
TITLE HRI human cDNA project  
JOURNAL Unpublished (2000)  
COMMENT Contact: Takao Isogai  
Genomics Laboratory  
Helix Research Institute  
1532-3 Yana, Kisarazu, Chiba 292-0812, Japan  
Tel: 81-438-52-3975

Fax: 81-438-52-3986  
Email: genomics@hri.co.jp  
HRI human cDNA project: 5'- & 3'-end one pass sequencing: Helix  
Research Institute; cDNA library construction: Department of  
Virology, Institute of Medical Science, University of Tokyo, and  
Helix Research Institute.  
Location/Qualifiers  
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Qy 661 ATCAGAGCCT 670  
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RESULT 7  
BG757448  
LOCUS

BG757448 829 bp mRNA linear EST 15-MAY-2001

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DEFINITION 602711092F1 NIH_MGC_48 Homo sapiens cDNA clone IMAGE:4851502 5',
            mRNA sequence.
ACCESSION  BG757448
VERSION     BG757448.1 GI:14068101
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SOURCE      Homo sapiens (human)
ORGANISM    Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE   1 (bases 1 to 829)
            NIH-MGC http://mgs.nci.nih.gov/.
AUTHORS     National Institutes of Health, Mammalian Gene Collection (MGC)
TITLE       Unpublished (1999)
JOURNAL     Contact: Robert Strausberg, Ph.D.
COMMENT      Email: cgabbs-remail.nih.gov
            Tissue Procurement: Louis M. Staudt, M.D., Ph.D.
            CDNA Library Preparation: Ling Hong/Rubin Laboratory
            CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
            DNA Sequencing by: Incyte Genomics, Inc.
            Clone distribution: MGC clone distribution information can be
            found through the I.M.A.G.E. Consortium/LLNL at:
            http://image.llnl.gov
            Plate: LLCM1698 row: 9 column: 23
            High quality sequence stop: 801.
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            /notes="Organ: B-cells; Vector: pOTB7; Site 1: XhoI;
            Site 2: EcoRI; CDNA made by oligo-dT priming.
            Directionally cloned into EcoRI/XhoI sites using the
            following 5' adaptor: GGCACGAG(G). Size-selected >500bp
            for average insert size 1.8kb. Library constructed by Ling
            Hong in the laboratory of Gerald M. Rubin (University of
            California, Berkeley) using ZAP-cDNA synthesis kit
            (Stratagene) and Superscript II RT (Life Technologies).
            Note: this is a NIH_MGC Library."
ORIGIN
Query Match 35.6%; Score 634.2; DB 4; Length 829;
Best Local Similarity 96.1%; Pred. No. 1.1e-159;
Matches 672; Conservative 0; Mismatches 23; Indels 4; Gaps 2;

QY 427 AGCATTCTTTATGAAGACTGGTCTTTTGATGGATGAAGAAAGGTCCAGTATGCTTCCT 486
    |||
Db 133 AGTACTTTTATGAAGACTGGTCTTTTGATGGATGAAGAAAGGTCCAGTATGCTTCCT 192

QY 487 ACCATGGCAGCAGGTCTGAATCCATCTCTTTGCGATTAACTCGACAAAGGATTGG 546
    |||
Db 193 ACCATGGCAGCAGGTCCGAACCTCATACTCTTTGCGATTAACTTGAACAACAGGATTGG 252

QY 547 AACGGGAGAGTAAGTTTGCTCCACCGTTTCAGACCTCTTAAGAGGTCAACGAGAAC 606
    |||
Db 253 AACGGGAGAGTAAGTTTGCTCCACCGTTTCAGACCTCTTAAGAGGTCAACGAGAAAT 312

QY 607 GTGACCTCTCTTGTGAAGAGGTCCACGCAAGGAGTGAGCAGCTGTTTCAGGGAGATCACA 666
    |||
Db 313 GTGA---CCTTGCTGAAGAGGTCAACCAAGGAGTGAGCAGCTGTTTCAGGAGATCACA 369

QY 667 GCCTCTCTCCGCTCTCCATCTCATCAAACTTGAACAGGAGACCGACCCCTTGCCTGTC 726
    |||
Db 370 GCCTCTCTCCATCTCTCCATCTCATCAAACTTGAACAGGAGACCGACCCCTTGCCTGTC 429

QY 727 GTGTCAGGAGATGTCAGTCTGATGCCAAATGCAAAAGAGCGGAGAGAGAAAGAA 786
    |||
Db 430 GTGTCCAGGAATGTGAGTCTGATGCCAAATGCAAAAGAGCGGAGAGAGAAAGAA 489

QY 787 GTGACCAACATAATCTCATTTGATGAGGAAGATGAGCAGAACTCTGGGGAGCTGTTT 846
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Db 490 GTGACCAACATTTATCTCATTTGATGAGGAAGATGAGCAGAACTCTGGGACATGTTT 549
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QY 847 AAAAAACACCTGGGGCAGGGAGAGCTCAGAGGACAACTCCGACCGTCTCTTGTCAAT 906
    |||
Db 550 AAAAAACACCTGGGGCAGGGAGAGCTCAGAGGACAACTCCGACCGTCTCTTGTCAAT 609
    |||
QY 907 ATCATGTCCGCTTTGAAAGCCCTTCGGCCCTAACTCCAATGGAAGTCAGAGCAGAAC 966
    |||
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Db 670 TCGTGAATAATTTGATTCCTCTCTTTTGAACCGGGAGTTTGGGTACCAGAAGCTTGTATG 729
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QY 1027 AAAAGCATCGATGATGAAGATGTGATGAAAACGAGATGACGTGTATGG-AACTCATC 1085
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Db 730 AAAAGCATCGATGATGAAGATGTGATGAAAACGAGATGACGTGTATGAAAACCTCATC 789
    |||
QY 1086 AGAAGGAGACACAGGGGCCACTCGGAGTCGCCCGAGAA 1124
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Db 790 AGGACGAAGCACATGTGGCACTCAGAGTCGGGGCGAGAA 828
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RESULT 8
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LOCUS 602712923F1 NIH_MGC_48 Homo sapiens cDNA clone IMAGE:4853188 5',
DEFINITION mRNA sequence.
ACCESSION  BG758677
VERSION     BG758677.1 GI:14069330
KEYWORDS    EST.
SOURCE      Homo sapiens (human)
ORGANISM    Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE   1 (bases 1 to 835)
            NIH-MGC http://mgs.nci.nih.gov/.
AUTHORS     National Institutes of Health, Mammalian Gene Collection (MGC)
TITLE       Unpublished (1999)
JOURNAL     Contact: Robert Strausberg, Ph.D.
COMMENT      Email: cgabbs-remail.nih.gov
            Tissue Procurement: Louis M. Staudt, M.D., Ph.D.
            CDNA Library Preparation: Ling Hong/Rubin Laboratory
            CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
            DNA Sequencing by: Incyte Genomics, Inc.
            Clone distribution: MGC clone distribution information can be
            found through the I.M.A.G.E. Consortium/LLNL at:
            http://image.llnl.gov
            Plate: LLCM1698 row: n column: 05
            High quality sequence stop: 819.
            Location/Qualifiers
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            /tissue_type="primary B-cells from tonsils (cell line)"
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            /clone_lib="NIH_MGC_48"
            /notes="Organ: B-cells; Vector: pOTB7; Site 1: XhoI;
            Site 2: EcoRI; CDNA made by oligo-dT priming.
            Directionally cloned into EcoRI/XhoI sites using the
            following 5' adaptor: GGCACGAG(G). Size-selected >500bp
            for average insert size 1.8kb. Library constructed by Ling
            Hong in the laboratory of Gerald M. Rubin (University of
            California, Berkeley) using ZAP-cDNA synthesis kit
            (Stratagene) and Superscript II RT (Life Technologies).
            Note: this is a NIH_MGC Library."
ORIGIN
Query Match 35.2%; Score 627.8; DB 4; Length 835;
Best Local Similarity 96.6%; Pred. No. 5.9e-158;

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Qy	487	ACATGTCAGAGGTCTGAACCTCATCTCTTTGCGATTAACTGCAACAAGGATTG							546
Db	91	ACATGTCAGAGGTCTGCAACTCCATCTCTTTGCGATTAACTGCAACAAGGATTG							147
Qy	547	AACGGGAGAGTAACTTTGCTCCACCGTTTCAGACCTCTTAAGGAGTCAAGCGAAC							606
Db	148	AACGGGAGAGTAACTTTGCTCCACCGTTTCAGACCTCTTAAGGAGTCAAGCGAAC							207
Qy	607	GTGACCTCTTGTGAGAGGAGTCCACCAAGGAGTGAAGAGTCTTTCAGGGAGATCACA							666
Db	208	GTGA---CCTTGTGAGAGGAGTCCACCAAGGAGTGAAGAGTCTTTCAGGGAGATCACA							264
Qy	667	GCCTCTCTGCGTCTCATCTCTCAAACTGAAAGAGACCGACCCCTTGCCTGTC							726
Db	265	GCCTCTCTGCGTCTCATCTCTCAAACTGAAAGAGACCGACCCCTTGCCTGTC							324
Qy	727	GTGTCAGGAGTGTGAGTCTGATGCCAAATGCAAAAGGAGCGGAAGAAAGAA							786
Db	325	GTGTCAGGAGTGTGAGTCTGATGCCAAATGCAAAAGGAGCGGAAGAAAGAA							384
Qy	787	GTGACCAACATAATCTCATTTGATGATGAGGAGATGAGCAGAACTCTGGGGAGCTGTT							846
Db	385	GTGACCAACATAATCTCATTTGATGATGAGGAGATGAGCAGAACTCTGGGGAGCTGTT							444
Qy	847	AAAAAGACACTGGGGAGGGAGAGCTCAGAGACAACTCCGACCCCTCTCTGTCAAT							906
Db	445	AAAAAGACACTGGGGAGGGAGAGCTCAGAGACAACTCCGACCCCTCTCTGTCAAT							504
Qy	907	ATCATGTCCGCTTTGAAAGCCCTTCGGGCTAACTCCAAATGGAAGTTCAGAGCAGAAC							966
Db	505	ATCATGTCCGCTTTGAAAGCCCTTCGGGCTAACTCCAAATGGAAGTTCAGAGCAGAAC							564
Qy	967	TCATGAAAAATTGATTCCTCTGTTTGAACGGGAGTTTGGGTACCAAGAGCTTGATGTG							1026
Db	565	TCGTGAAAAATTGATTCCTCTGTTTGAACAGGAGTTTGGGTACCAAGAGCTTGATGTG							624
Qy	1027	AAAGCATCGATGATGAAGATGAGATGAAACGAAGATGACGTGTATGAAACTCATC							1085
Db	625	AAAGCATCGATGATGAAGATGAGATGAAACGAAGATGACGTGTATGAAACTCATC							684
Qy	1086	AGGAGAACACAGGGGCGCACTCGGAGTCGCCCGAGAA	1124						
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RESULT 9  
BO057616  
LOCUS  
DEFINITION  
AGENCOURT\_7047416 NIH\_MGC\_99 Homo sapiens cdna clone IMAGE:5813279  
5', mRNA sequence.  
ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
1 (bases 1 to 1022)  
NIH-MGC <http://mgi.nci.nih.gov/>.  
National Institutes of Health, Mammalian Gene Collection (MGC)  
Unpublished (1999)  
Contact: Robert Strausberg, Ph.D.  
Email: [cgabs-re@mail.nih.gov](mailto:cgabs-re@mail.nih.gov)  
Tissue Procurement: Lou Staudt  
CDNA Library Preparation: Rubin Laboratory  
CDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
DNA Sequencing by: Agencourt Bioscience Corporation

Clone distribution: MGC clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: <http://image.llnl.gov>  
Plate: L1CM2065 row: a column: 24  
High quality sequence stop: 641.  
Location/Qualifiers  
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/clone="IMAGE:5813279"  
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/clone\_lib="NIH MGC 99"  
/notes="Organ: lymph; Vector: pOTB7; Site 1: XhoI; Site 2: EcoRI; cDNA made by oligo-dT priming. Directionally cloned into EcoRI/XhoI sites using the following 5' adaptor: GGACAG(G). Size-selected >500bp for average insert size 1.8kb. Library constructed by Ling Hong in the laboratory of Gerald M. Rubin (University of California, Berkeley) using ZAP-cDNA synthesis kit (Stratagene) and Superscript II RT (Life Technologies). Note: this is a NIH\_MGC Library."  
ORIGIN  
Query Match 35.2%; Score 627.8; DB 5; Length 1022;  
Best Local Similarity 96.3%; Pred. No. 6.3e-158;  
Matches 654; Conservative 0; Mismatches 22; Indels 3; Gaps 1;  
Qy 427 AGCACTTTTATGAAGACTGGTCTTTTGTGATGATGAAGAAAGGTCCAGTATGCTTCCT 486  
Db 70 AGTACTTTTATGAAGACTGGTCTTTTGTGATGATGAAGAGGTCCAGTATGCTTCCT 129  
Qy 487 ACCATGTCAGAGGTCTGAACCTCATCTCTTTGCGATTAACTGCAACAAGGATTG 546  
Db 130 ACCATGTCAGAGGTCTGAACCTCATCTCTTTGCGATTAACTGCAACAAGGATTG 189  
Qy 547 AACGGGAGAGTAACTTTGCTCCACCGTTTCAGACCTCTTAAGGAGTCAAGCGAAC 606  
Db 190 AACGGGAGAGTAACTTTGCTCCACCGTTTCAGACCTCTTAAGGAGTCAAGCGAAC 249  
Qy 607 GTGACCTCTTGTGAGGAGTCCACCAAGGAGTGAAGAGTCTTTCAGGGAGATCACA 666  
Db 250 GTGA---CCTTGTGAGGAGTCCACCAAGGAGTGAAGAGTCTTTCAGGGAGATCACA 306  
Qy 667 GCCTCTCTGCGTCTCATCTCTCAAACTGAAAGAGACCGACCCCTTGCCTGTC 726  
Db 307 GCCTCTCTGCGTCTCATCTCTCAAACTGAAAGAGACCGACCCCTTGCCTGTC 366  
Qy 727 GTGTCCAGGAATGTGAGTCTGATGCCAAATGCAAAAGGAGCGGAAGAAAGAA 786  
Db 367 GTGTCCAGGAATGTGAGTCTGATGCCAAATGCAAAAGGAGCGGAAGAAAGAA 426  
Qy 787 GTGACCAACATAATCTCATTTGATGATGAGGAGATGAGCAGAACTCTGGGGAGCTGTT 846  
Db 427 GTGACCAACATAATCTCATTTGATGATGAGGAGATGAGCAGAACTCTGGGGAGCTGTT 486  
Qy 847 AAAAAAGACACTGGGGAGGGAGAGCTCAGAGACAACTCCGACCCCTCTCTGTCAAT 906  
Db 487 AAAAAAGACACTGGGGAGGGAGAGCTCAGAGACAACTCCGACCCCTCTCTGTCAAT 546  
Qy 907 ATCATGTCCGCTTTGAAAGCCCTTCGGGCTAACTCCAAATGGAAGTTCAGAGCAGAAC 966  
Db 547 ATCATGTCCGCTTTGAAAGCCCTTCGGGCTAACTCCAAATGGAAGTTCAGAGCAGAAC 606  
Qy 967 TCATGAAAAATTGATTCCTCTGTTTGAACGGGAGTTTGGGTACCAAGAGCTTGATGTG 1026  
Db 607 TCGTGAATAATGATTCCTCTGTTTGAACAGGAGTTTGGGGACCAAGAGCTTGATGTG 666  
Qy 1027 AAAAGCATCGATGATGAAGATGAGATGAAACGAAGATGACGTGTATGAAACTCATCA 1086  
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FEATURES  
source





of Gerald M. Rubin (University of California, Berkeley)  
using ZAP-cDNA synthesis kit (Stratagene) and Superscript  
II RT (Life Technologies). Note: this is a NIH\_MGC  
Library."

ORIGIN	Query Match 35.0%; Score 624.4; DB 5; Length 892; Best Local Similarity 92.4%; Pred. No. 5e-157; Matches 680; Conservative 0; Mismatches 51; Indels 5; Gaps 2;
Qy	427 AGCACTTTTATGAAGACTGGTCTTTTGTGATGATGAAGAAAGTCCAGTATGCTTCT 486
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Qy	487 ACCATGGCAGAGGTCTGAATCCATCTCTTTGGATTAAATCATGCAACAAGGATTTG 546
Db	158 ACCATGGCAGAGGTCCGAATCCATCTCTTTGGATTAAATCATGCAACAAGGATTTG 217
Qy	547 AACGGGACAGTAACTTTGCTCCACCTTTTCAGACCTTTAAAGGAGTCAACGGAGAAC 606
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Qy	607 GTGACCTCTTGTGTAAGAGTCCACGCAAGAGTGACGAGCTGTTTCAGGGAGATCACA 666
Db	278 GTGA---CCTTGTGTAAGAGTCCACGCAAGAGTGACGAGCTGTTTCAGGGAGATCACA 334
Qy	667 GCCTCTCTGCGCTCTCCATCTCTCATCAAACTGAAACAGGAGACCGACCCCTTGCTGTC 726
Db	335 GCCTCTCTGCGCTCTCCATCTCTCATCAAACTGAAACAGGAGACCGACCCCTTGCCGTC 394
Qy	727 GTGTCCAGGAATGTCAGTGTGATGCCAAATGCAAAAGGAGCGGAAGAAAGAA 786
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Qy	787 GTGACCAACAATCTCTATTTGATGATCAGGAAGATGACGAGAACTCTGGGAGCTGTTT 846
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Qy	847 AAAAAGACACCTGGGGGAGGAGAGCTCAGAGGACAACTCCGACCGCTCTCTGTCTCAAT 906
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Qy	1027 AAAAGCATCGATGATGAAGATGTGGATGAAACCGAAGATGACGTGTATGGAACTCATCA 1086
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Qy	1087 GGAA--GGAAGCAGAGGGGCACTCGGAGTCCGCCGAGAACCACTGGAAGGGAAACACT 1144
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Qy	1145 GCCTCTCCACATGCA 1160
Db	815 GTCACCCCGCTGGA 830
RESULT 12	
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DEFINITION	UI-E-CK1-afk-h-18-0-UI.r1 UI-E-CK1 Homo sapiens cDNA clone
ACCESSION	UI-E-CK1-afk-h-18-0-UI 5', mRNA sequence.
VERSION	BM692924
KEYWORDS	BM692924.1 GI:19006182
SOURCE	EST.
ORGANISM	Homo sapiens (human)

REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
AUTHORS	1 (bases 1 to 609)
TITLE	Bonaldo,M.F., Lennon,G. and Soares,M.B. Normalization and subtraction: two approaches to facilitate gene discovery
JOURNAL	Genome Res. 6 (9), 791-806 (1996)
MEDLINE	97044477
PUBMED	8889548
COMMENT	Contact: Soares, MB Coordinated Laboratory for Computational Genomics University of Iowa 375 Newton Road, 4156 MEBRP, Iowa City, IA 52242, USA Tel: 319 335 8250 Fax: 319 335 9565 Email: bento-soares@uiowa.edu Tissue Procurement: Dr. Gregg Hageman cDNA Library preparation: Dr. M. Bento Soares, University of Iowa cDNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa DNA Sequencing by: Dr. M. Bento Soares, University of Iowa Clone Distribution: Researchers may obtain clones from Research Genetics (www.resgen.com). Seq primer: M13 Reverse.
FEATURES	Location/Qualifiers
source	1..609 /organism="Homo sapiens" /mol_type="mRNA" /db_xref="taxon:9606" /clone="UI-E-CK1-afk-h-18-0-UI" /tissue_type="Retina Foveal and Macular" /dev_stage="adult" /lab_host="DH10B (Life Technologies) (T1 phage resistant)" /clone_lib="UI-E-CK1" /note="Organ: eye; Vector: pTVT3-Pac (Pharmacia) with a modified polylinker; Site 1: EcoR I; Site 2: Not I; UI-E-CK1 is a normalized cDNA library containing the following tissue(s): Retina Foveal and Macular. The library was constructed according to Bonaldo, Lennon and Soares, Genome Research, 6:791-806, 1996. First strand cDNA synthesis was primed with an oligo-dT primer containing a Not I site. Double stranded cDNA was ligated to an EcoR I adaptor, digested with Not I, and cloned directionally into pTVT3-Pac vector. The oligonucleotide used to prime the synthesis of first-strand cDNA contains a library tag sequence that is located between the Not I site and the (dT)18 tail. The sequence tag for this library is GTCC. This library was created for the program, Gene Discovery in the Visual System, supported by National Eye Institute (NEI)."
ORIGIN	
Query Match 33.6%; Score 598; DB 4; Length 609; Best Local Similarity 99.8%; Pred. No. 6.1e-150; Matches 609; Conservative 0; Mismatches 0; Indels 1; Gaps 1;	
Qy	1100 GGGGCCACTCGGAGTCGCCGAGAACCACTGGAAGGGAACACCTGCTCTCCAGATGC 1159
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 Db 301 TTAGTGAACCTGCCAGGCCACTGTGGCCATGATGAACAGGAAGGATGAGCTGGAGGAG 360  
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 Qy 1580 TCAGGCGCTGGCCAGCACTATCTTTGTCTATTTGTGAGGAGATTCTAACCCCAAGTGA 1639  
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 Qy 1700 GCTCTCTGGAT 1709  
 Db 600 GCTCTCTGGAT 609

## RESULT 13

BG684262

LOCUS

DEFINITION 813 bp mRNA linear EST 01-MAY-2001  
 602635896F1 NIH\_MGC\_48 Homo sapiens cDNA clone IMAGE:4763783 5',  
 mRNA sequence.

ACCESSION

BG684262

VERSION

1

KEYWORDS

EST.

SOURCE

Homo sapiens

ORGANISM

Homo sapiens

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

BG684262 813 bp mRNA linear EST 01-MAY-2001  
 602635896F1 NIH\_MGC\_48 Homo sapiens cDNA clone IMAGE:4763783 5',  
 mRNA sequence.  
 BG684262 1 GI:13915659  
 EST.  
 Homo sapiens (human)  
 Homo sapiens  
 Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
 1 (bases 1 to 813)  
 NIH-MGC http://mgs.nci.nih.gov/  
 National Institutes of Health, Mammalian Gene Collection (MGC)  
 Unpublished (1999)  
 Contact: Robert Strausberg, Ph.D.  
 Email: cgaabs@mail.nih.gov  
 Tissue Procurement: Louis M. Staudt, M.D., Ph.D.  
 cDNA Library Preparation: Ling Hong/Rubin Laboratory  
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
 DNA Sequencing by: Incyte Genomics, Inc.  
 Clone distribution: MGC clone distribution information can be  
 found through the I.M.A.G.E. Consortium/LLNL at:  
 http://image.llnl.gov  
 Plate: LLCMI619 row: p column: 24  
 High quality sequence stop: 795.  
 Location/Qualifiers

## FEATURES

source

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 /notes="Organ: B-cells; Vector: pOTB7; Site:1: XhoI;  
 Site:2: EcoRI; cDNA made by oligo-dT priming.  
 Directionally cloned into EcoRI/XhoI sites using the  
 following 5' adaptor: GGCAGAG(G). Size-selected >500bp  
 for average insert size 1.8kb. Library constructed by Ling  
 Hong in the laboratory of Gerald M. Rubin (University of  
 California, Berkeley) using ZAP-cDNA synthesis kit  
 (Stratagene) and Superscript II RT (Life Technologies).  
 Note: this is a NIH\_MGC Library."

ORIGIN

Query Match 32.3%; Score 574.8; DB 4; Length 813;  
 Best Local Similarity 96.3%; Pred. No. 1.2e-143;  
 Matches 653; Conservative 0; Mismatches 17; Indels 8; Gaps 6;  
 QY 427 AGCACATTTTATGAAGACTGGTCTTTTGTGATGATGAAGAAAGTCCAGTATGCTTCCT 486  
 DB 142 AGTACTTTTATGAAGACTGGTCTTTTGTGATGATGAAGAAAGTCCAGTATGCTTCCT 200  
 QY 487 ACCATGGCAGCAGGTCTGAACCTCCATCTCTTTTGGGATTAAACATCGACAAACAGGATTTG 546  
 DB 201 ACCATGGCAGCAGGTCCGAATCCATCTCTTTGGGATTAAACATTCACAAACAAAGATTG 260  
 QY 547 AACGGGCAGAGTAAGTTTGTCTCCACCGTTTCCAGACCTCTTAAAGAGTCAACGAGAAC 606  
 DB 261 AACGGGCAGAGTAAGTTTGTCTCCACCGTTTCCAGACCTCTTAAAGAGTCAACGAGAAC 320  
 QY 607 GTGACCTCTCTGTGAAGGAGTCCACGCAAGAGTCCAGCAGCTTTTCCAGGGAGATCACA 666  
 DB 321 GTGA----CTTGTCTGAAGAGTCCACGCAAGAGTCCAGCAGCTTTTCCAGGGAGATCACA 377  
 QY 667 GCTCTCTCTGCGCTCTCCATCTCTCATCAAAACCTGAAACAGACCAACCCCTTCCCTGCTC 736  
 DB 378 GCTCTCTCTGCGCTCTCCATCTCTCATCAAAACCTGAAACAGACCAACCCCTTCCCTGCTC 437  
 QY 727 GTGTCAGGAATGTCTAGTCTCTGATGCCAAATGCCAAAGAGCGGGAAGAAAGAA 786  
 DB 438 GTGTCAGGAATGTCTAGTCTCTGATGCCAAATGCCAAAGAGCGGGAAGAAAGAAAGCAA 497  
 QY 787 GTGACCAACATATCTCTTGTGATGATGAGGAAGATGACAGAACTCTCGGGACGTGTTT 846  
 DB 498 GTGACCAACATATCTCTTGTGATGATGAGGAAGATGACAGAACTCTCGGGACGTGTTT 557  
 QY 847 AAAAAA-CACCTGGGCGCAGGGAGAGCTCAGAGGACAACTCCGACCGCTCTCTCTGCTCAA 905  
 DB 558 AAAAAACACCTGGGCGCAGGGAGAGCTCAGAGGACAACTCCGACCGCTCTCTCTGCTCAA 616  
 QY 906 TATCATGTGCGCTTTTGAAGCCCTTCCGGCTTAATCAATGGAAGTCCAGAGACAA 965  
 DB 617 TATCATGTGCGCTTTGACAGCCCTTCCGGGCCAACTCCAATGGAAGTCCAGAGACAA 676  
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 DB 677 CTCGTGAAAAATTTGGATTCCCTGCTCTTGAACAGGGAG-TTGGGTACCAAGACTTGTATG 735  
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 DB 736 TGAAGAGCATCGATGATGAAGATGTGGATGAAACCAAGATGACGTGTATGGAACATCAT 795  
 QY 1085 CAGGAAGGAGCACAGGG 1102  
 DB 796 CAGGACGGAAGCACAGGG 813

## RESULT 14

BG759681

LOCUS

DEFINITION

602713374F1 NIH\_MGC\_48 Homo sapiens cDNA clone IMAGE:4853346 5',

mRNA sequence.

BG759681

VERSION

1

KEYWORDS

EST.

SOURCE

Homo sapiens

ORGANISM

Homo sapiens

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

BG759681 819 bp mRNA linear EST 15-MAY-2001  
 602713374F1 NIH\_MGC\_48 Homo sapiens cDNA clone IMAGE:4853346 5',  
 mRNA sequence.  
 BG759681 1 GI:14070334  
 EST.  
 Homo sapiens (human)  
 Homo sapiens  
 Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
 1 (bases 1 to 819)  
 NIH-MGC http://mgs.nci.nih.gov/  
 National Institutes of Health, Mammalian Gene Collection (MGC)  
 Unpublished (1999)  
 Contact: Robert Strausberg, Ph.D.  
 Email: cgaabs@mail.nih.gov  
 Tissue Procurement: Louis M. Staudt, M.D., Ph.D.

cDNA Library Preparation: Ling Hong/Rubin Laboratory  
cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
DNA Sequencing by: Incyte Genomics, Inc.  
Clone distribution: MGC clone distribution information can be  
found through the I.M.A.G.E. Consortium/LLNL at:  
<http://image.llnl.gov>  
Place: LUCM1699 row: d column: 19  
High quality sequence stop: 791.  
Location/Qualifiers

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/clone\_lib="NIH MGC 48"  
/note="Organ: B-cells; Vector: pOTB7; Site1: XhoI;  
Site2: EcoRI; cDNA made by oligo-dT priming.  
Directionally cloned into EcoRI/XhoI sites using the  
following 5' adaptor: GGCACGAG(G). Size-selected >500bp  
for average insert size 1.8kb. Library constructed by Ling  
Hong in the laboratory of Gerald M. Rubin (University of  
California, Berkeley) using ZAP-cDNA synthesis kit  
(Stratagene) and Superscript II RT (Life Technologies).  
Note: this is a NIH\_MGC Library."

## ORIGIN

Query Match 30.0%; Score 535; DB 4; Length 819;  
Best Local Similarity 95.4%; Pred. No. 7e-133;  
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Db |||  
Qy 487 ACCATGGCAGAGGTCTGAATCCATACTCTTTTGGATTATCATCGACAAACAAGATTG 546  
Db |||  
Qy 237 ACCATGGCAGAGGTCCGAATCCATACTCTTTTGGATTATCATCGACAAACAAGATTG 296  
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Qy 607 GTGACCTCTTCTGAAGAGTCCACGCAAGAGTGAAGAGTGAAGAGTGAAGAGTGAAG 666  
Db |||  
Qy 357 GTGA---CCTTGTGAAGAGTCCACGCAAGAGTGAAGAGTGAAGAGTGAAGAGTGAAG 413  
Qy 667 GCCTCTCTGCGCTCTCCATCTCATCAAACTGAACAGGAGACGACCCCTTTCCTGTGTC 726  
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Qy 414 GCCTCTCTGCGCTCTCCATCTCATCAAACTGAACAGGAGACGACCCCTTTCCTGTGTC 473  
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Db |||  
Qy 787 GTGACCAACATAATCTCATTTTGTGATGATGAGGAAGTGAAGAGTCTTGGGGAGCTGTT 846  
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Qy 534 GTGACCAACATAATCTCATTTTGTGATGATGAGGAAGTGAAGAGTCTTGGGGAGCTGTT 593  
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Qy 594 AAAAAAGACACCTGGGGAGGGAGAGCTCAGAGGACAACTCCAGCCCTCTCTGTCAA 653  
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Qy 654 TATCATGTCGCCCTTTGAAG-CCCTTCGGCCCAACTCCCAATGAAGTCAAGACGACGAA 712  
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Db 773 TGTGAAGCATCGATGATGAAGATGTGGGATCAAAACGAAGATTA 819  
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RESULT 15  
BQ184171/c  
LOCUS

DEFINITION  
UI-E-EJ1-ajs-e-23-0-UI.s1 UI-E-EJ1 Homo sapiens cDNA clone  
UI-E-EJ1-ajs-e-23-0-UI 3', mRNA sequence.

ACCESSION  
BQ184171  
VERSION  
BQ184171.1  
KEYWORDS  
EST.

SOURCE  
Homo sapiens (human)

ORGANISM  
Homo sapiens  
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE  
AUTHORS  
TITLE

1 (bases 1 to 626)  
Bonaldo, M.F., Lennon, G. and Soares, M.B.  
Normalization and subtraction: two approaches to facilitate gene  
discovery

JOURNAL  
MEDLINE  
PUBMED  
COMMENT

Genome Res. 6 (9), 791-806 (1996)  
97044477  
8889548  
Contact: Soares, MB  
Coordinated Laboratory for Computational Genomics  
University of Iowa  
375 Newton Road, 4156 MEBRF, Iowa City, IA 52242, USA

Tel: 319 335 8250  
Fax: 319 335 9565

Email: bento-soares@uiowa.edu  
Tissue Procurement: Dr. Gregg Hageman  
cDNA Library preparation: Dr. M. Bento Soares, University of Iowa  
cDNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa

DNA Sequencing by: Dr. M. Bento Soares, University of Iowa  
Clone Distribution: Researchers may obtain clones from Research  
Genetics (www.resgen.com).

Seq primer: M13 FORWARD  
POLYA=Yes.

FEATURES  
source

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/tissue="fetal eyes, lens, eye anterior segment,  
optic nerve, retina, Retina Foveal and Macular, RPE and  
Choroid"

/dev\_stage="fetal and adult"  
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/clone\_lib="UI-E-EJ1"

/note="Organ: eye; Vector: pT7T3-Pac (Pharmacia) with a  
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UI-E-EJ1 is a subcloned cDNA library constructed  
according to Bonaldo, Lennon and Soares, Genome Research,  
6:791-806, 1996. First strand cDNA synthesis was primed  
with an oligo-dT primer containing a Not I site. Double  
stranded cDNA was ligated to an EcoR I adaptor, digested  
with Not I, and cloned directionally into pT7T3-Pac  
vector. The oligonucleotide used to prime the synthesis of  
first-strand cDNA contains a library tag sequence that is  
located between the Not I site and the (dT)18 tail. The  
sequence tags for this library are: fetal eyes,  
AGATCAAGA; lens, CGATTAGCA; eye anterior segment,  
AATGCCGAT; optic nerve, CCATTAGTG; retina, CCGCG; Retina  
Foveal and Macular, GTCC; RPE and Choroid, ACCTA. This  
library was created for the program, Gene Discovery in the  
Visual System, supported by National Eye Institute (NEI).

TAG TISSUE=human retina  
TAG\_LIB=UI-E-EJ1  
TAG\_SEQ=CCGCG"

ORIGIN

Query Match 30.0%; Score 534.6; DB 5; Length 626;

Host Local Similarity 99.1%; Pred. No. 8.3e-133;		Matches 537; Conservative 0; Mismatches 5; Indels 0; Gaps 0;	
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QY	425	TGAGCACTTTTATGAAGACTGCTCTTTTGTGTGATGATGAAGAAAGTCCAGTATGCTTC	484
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QY	665	CAGCCTCCTCTGCGGTCTCCATCCTCATCAACCTGACAGGAGACCGACCCCTTGCCCTG	724
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QY	725	TCGTGTCCAGGAATGTCAAGTGTGATGCCCAAAATGCAAAAAGGAGCGGAAGAAAAAGA	784
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Db		4 AA 3	

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GenCore version 5.1.6  
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OM nucleic - nucleic search, using sw model

Run on: October 25, 2005, 21:11:53 ; Search time 1482 Seconds  
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Post-processing: Minimum Match 0%  
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Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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1	275.8	15.5	280	9	US-09-563-817-201
2	247	13.9	454	24	Sequence 201, App
C 3	227.8	12.8	542	17	Sequence 214, App
C 4	224.8	12.6	284	17	Sequence 8542, App
5	207.6	11.6	463	24	Sequence 22242, A

6	206.8	11.6	465	26	US-11-060-756-3234	Sequence 3234, App
7	206.8	11.6	465	26	US-11-060-756-7506	Sequence 7506, App
C 8	143.8	8.1	492	13	US-09-925-065A-383678	Sequence 383678, App
C 9	135.4	7.6	474	24	US-10-450-763-9353	Sequence 9353, App
10	128	7.2	500	17	US-10-029-386-8336	Sequence 8336, App
11	124	7.0	124	17	US-10-029-386-22036	Sequence 22036, App
C 12	119	6.7	911	24	US-10-450-763-215	Sequence 215, App
13	75	4.2	633	13	US-09-925-065A-681221	Sequence 681221, App
14	66.8	3.7	1796	26	US-11-097-143-11288	Sequence 11288, App
C 15	66.8	3.7	3796	26	US-11-097-143-11287	Sequence 11287, App
C 16	63.8	3.6	3931	19	US-10-276-774-948	Sequence 948, App
17	63	3.5	166536	24	US-10-981-277-35	Sequence 35, Appl
18	54.4	3.1	813	24	US-10-450-763-21166	Sequence 21166, App
C 19	54.4	3.1	79977	24	US-10-737-082-58	Sequence 58, Appl
C 20	54.4	3.1	79977	24	US-10-765-790-58	Sequence 58, Appl
21	54	3.0	464	10	US-09-918-995-15177	Sequence 15177, App
C 22	52.8	3.0	263	9	US-09-867-701-9696	Sequence 9696, App
23	52.8	3.0	476	10	US-09-854-867-118	Sequence 118, App
24	52.8	3.0	476	22	US-10-786-970A-118	Sequence 118, App
25	50	2.8	371	10	US-09-918-995-24284	Sequence 24284, App
C 26	50	2.8	446	9	US-09-880-107-1518	Sequence 1518, App
27	50	2.8	7093	16	US-10-101-510-527	Sequence 527, App
28	50	2.8	10126	18	US-10-242-355-1024	Sequence 1024, App
29	49.6	2.8	7140	22	US-10-211-028-3	Sequence 3, Appl
30	49.6	2.8	90597	22	US-10-211-028-1	Sequence 1, Appl
31	49.2	2.8	18535	15	US-10-079-854-385	Sequence 385, App
32	49.2	2.8	2463	14	US-10-027-632-102953	Sequence 102953, App
33	48	2.7	2463	18	US-10-027-632-102953	Sequence 102953, App
34	48	2.7	2927	24	US-10-450-763-24335	Sequence 24335, App
C 35	48	2.7	2927	24	US-10-450-763-24335	Sequence 24335, App
36	47.6	2.7	5430	24	US-11-097-143-14315	Sequence 14315, App
37	47.2	2.6	2227	26	US-11-097-143-14314	Sequence 14314, App
38	47.2	2.6	4602	23	US-09-925-065A-844196	Sequence 844196, App
C 39	46.6	2.5	473	13	US-10-027-632-276057	Sequence 276057, App
C 40	44.4	2.5	505	14	US-10-027-632-276057	Sequence 276057, App
C 41	44.4	2.5	505	18	US-10-437-963-53652	Sequence 53652, App
C 42	44	2.5	4216	20	US-10-437-963-53652	Sequence 53652, App
C 43	43.8	2.5	1599	20	US-10-437-963-2771	Sequence 2771, App
C 44	43.2	2.4	505	14	US-10-027-632-276059	Sequence 276059, App
C 45	43.2	2.4	505	18	US-10-027-632-276059	Sequence 276059, App

ALIGNMENTS

RESULT 1

US-09-563-817-201  
; Sequence 201, Application US/09563817  
; Patent No. US20020095031A1  
; GENERAL INFORMATION:  
; APPLICANT: Nehls, Michael C.  
; APPLICANT: Zambrowicz, Brian  
; APPLICANT: Sands, Arthur T.  
; TITLE OF INVENTION: No. US20020095031A1 Human Polynucleotides and the  
; FILE OF INVENTION: Polypeptides Encoded Thereby  
; FILE REFERENCE: LEX-0021-USA  
; CURRENT APPLICATION NUMBER: US/09/563.817  
; CURRENT FILING DATE: 2001-06-11  
; PRIOR APPLICATION NUMBER: US 60/132,343  
; PRIOR FILING DATE: 1999-05-04  
; NUMBER OF SEQ ID NOS: 1008  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 201  
; LENGTH: 280  
; TYPE: DNA  
; ORGANISM: homo sapiens  
; FEATURE:  
; NAME/KEY: misc\_feature  
; LOCATION: (1)..(280)  
; OTHER INFORMATION: n = A,T,C or G  
US-09-563-817-201

Query Match 15.5% Score 275.8 DB 9 Length 280



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QY	1399	ATTAGTGAATCTGCGCAGGCCACTGTGGCCATGATGAACAGGAAGGATGA	1448
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RESULT 6			
US-11-060-756-3234			
; Sequence 3234, Application US/11060756			
; Publication No. US20050221354A1			
; GENERAL INFORMATION:			
; APPLICANT: Wyeth			
; APPLICANT: Mounts, William Martin			
; TITLE OF INVENTION: Nucleic Acid Arrays for Monitoring Expression Profiles of Drug			
; TITLE OF INVENTION: Target Genes			
; FILE REFERENCE: AM101083 (031896-042000)			
; CURRENT APPLICATION NUMBER: US/11/060,756			
; CURRENT FILING DATE: 2005-02-18			
; NUMBER OF SEQ ID NOS: 303284			
; SOFTWARE: PatentIn version 3.2			
; SEQ ID NO 3234			
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; TYPE: DNA			
; ORGANISM: Homo sapiens			
US-11-060-756-3234			
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Db	219	CTGCAACATGCTCTCTGGCGCAGCGCTGAGGCTGAGCACTTTTTATGAAGACTGGTCTTT	278
QY	453	TGTGATGGATGAAGAAGGTCCAGTATGCTTCTTACCATGGCAGCAGGTCTGGAATCC	510
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US-11-060-756-7506			
; Sequence 7506, Application US/11060756			
; Publication No. US20050221354A1			
; GENERAL INFORMATION:			
; APPLICANT: Wyeth			
; APPLICANT: Mounts, William Martin			
; TITLE OF INVENTION: Nucleic Acid Arrays for Monitoring Expression Profiles of Drug			
; TITLE OF INVENTION: Target Genes			
; FILE REFERENCE: AM101083 (031896-042000)			
; CURRENT APPLICATION NUMBER: US/11/060,756			
; CURRENT FILING DATE: 2005-02-18			
; NUMBER OF SEQ ID NOS: 303284			
; SOFTWARE: PatentIn version 3.2			
; SEQ ID NO 7506			
; LENGTH: 465			
; TYPE: DNA			
; ORGANISM: Homo sapiens			
US-11-060-756-7506			

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Best Local Similarity 94.5%; Pred. No. 4.6e-53;
Matches 225; Conservative 0; Mismatches 12; Indels 1; Gaps 1;

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Db 159 CGTGGCTGGGCGCGCGCTGGCTGGCTGTGCGCTCAACGAACACTCCCTGGAGTGCTA 218
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QY 393 CTGACATGCTCTCTGGCGACCGCTGCAGGCTGAGCACTTTTATGAAGACTGCTTTT 452
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Db 219 CTGACATGCTCTCTGGCGACCGCTGCAGGCTGAGCACTTTTATGAAGACTGCTTTT 278
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## RESULT 8

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US-09-925-065A-383678/c
; Sequence 383678, Application US/09925065A
; Publication No. US20050228172A9
```

```
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
```

```
; TITLE OF INVENTION: Identification and Mapping of Single
; Nucleotide Polymorphisms in the Human Genome
; FILE REFERENCE: 108827.135
```

```
; CURRENT APPLICATION NUMBER: US/09/925, 065A
; CURRENT FILING DATE: 2001-08-08
```

```
; PRIOR APPLICATION NUMBER: US 60/243,096
; PRIOR FILING DATE: 2000-10-24
```

```
; PRIOR APPLICATION NUMBER: US 60/252,147
; PRIOR FILING DATE: 2000-11-20
```

```
; PRIOR APPLICATION NUMBER: US 60/250,092
; PRIOR FILING DATE: 2000-11-30
```

```
; PRIOR APPLICATION NUMBER: US 60/261,766
; PRIOR FILING DATE: 2001-01-16
```

```
; PRIOR APPLICATION NUMBER: US 60/289,846
; PRIOR FILING DATE: 2001-05-09
```

```
; NUMBER OF SEQ ID NOS: 957086
; SOFTWARE: FASTSEQ for Windows Version 4.0
```

```
; SEQ ID NO 383678
; LENGTH: 492
```

```
; TYPE: DNA
; ORGANISM: Homo sapiens
```

```
US-09-925-065A-383678
```

```
Query Match      8.1%; Score 143.8; DB 13; Length 492;
Best Local Similarity 98.1%; Pred. No. 1.9e-33;
Matches 156; Conservative 0; Mismatches 2; Indels 1; Gaps 1;
```

```
QY 1589 TGCCAGCTATCTTTGCTATTTGTGAGGAGATCTAACCCACGTCGAGAACCATGTGGT 1648
    |||||
Db 158 TTGGCAGCTATCTTTGCTATTTGTGAGGAGATCTAACCCACGTCGAGAACCATGTGGT 99
    |||||
```

```
QY 1649 GGAGAAATGGAGGAGAGAGAAATCCAAAGTTTCTCTGATAGTCTCATTTGAGCTCTCTGGA 1708
    |||||
Db 98 GGAGAAATGGAGGAGAGAGAAAT-CAACAGTTCTCTGATAGTCTCATTTGAGCTCTCTGGA 40
    |||||
```

```
QY 1709 TCCAGTCTTTTCCCTGAAGCTGTGTTTCTCTGGACTTTTC 1747
    |||||
Db 39 TCCAGTCTTTTCCCTGAAGCTGTGTTTCTCTGGACTTTTC 1
    |||||
```

## RESULT 9

```
US-10-450-763-9353/c
; Sequence 9353, Application US/10450763
; Publication No. US20050196754A1
```

```
; GENERAL INFORMATION:
; APPLICANT: Penn, Sharron G.
```

```
; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
; THE ANALYSIS OF EXPRESSION ANALYSIS TWO
```

```
; FILE REFERENCE: AEOMICA-X-2
; CURRENT APPLICATION NUMBER: US/10/029,386
; CURRENT FILING DATE: 2001-12-20
```

```
; NUMBER OF SEQ ID NOS: 34288
; SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
; SEQ ID NO 8336
; LENGTH: 500
```

```
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: MAP TO CHR16.3
```

```
; OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 1.4
; OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 1.4
; OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 1.2
```

```
; APPLICANT: Hyseq, Inc
; TITLE OF INVENTION: NOVEL NUCLEIC ACIDS AND POLYPEPTIDES
; FILE REFERENCE: 790CIP3/US
; CURRENT APPLICATION NUMBER: US/10/450,763
; CURRENT FILING DATE: 2003-06-11
; PRIOR APPLICATION NUMBER: PCT/US01/08631
; PRIOR FILING DATE: 2001-03-30
; PRIOR APPLICATION NUMBER: 09/540,217
; PRIOR FILING DATE: 2000-03-31
; PRIOR APPLICATION NUMBER: 09/649,167
; PRIOR FILING DATE: 2000-08-23
; NUMBER OF SEQ ID NOS: 60736
; SOFTWARE: Custom
; SEQ ID NO 9353
; LENGTH: 474
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: SIMILAR
; LOCATION: (369)..(133)
; OTHER INFORMATION: 67% homologous to Homo sapiens Partial sequence of the clone
; OTHER INFORMATION: 9 protein, accession number W42400, Smith-Waterman Score=227.
US-10-450-763-9353
```

```
Query Match      7.6%; Score 135.4; DB 24; Length 474;
Best Local Similarity 72.6%; Pred. No. 7.5e-31;
Matches 175; Conservative 0; Mismatches 66; Indels 0; Gaps 0;
```

```
QY 81 CTTTGAGGAGGAGAGGAGATTGCCCTCGGATTCGACAGCAGGCTCACCTGTCTGTGTGC 140
    |||||
Db 367 CTCTGCCAGAGCAGAGAAAGTTCTGACCACTGGCAGAGGTCACCTGTCTGTGTGC 308
    |||||
```

```
QY 141 CCAGTTTGAAGCGCTCTCGCATGGCTTGAAGAGAGTTCGAGGATTCGACTCAGC 200
    |||||
Db 307 CCAGTTTGAAGCGCTCTCGCATGGCTTGAAGAGAGTTCGAGGATTCGACTCAGC 248
    |||||
```

```
QY 201 GCAGCGATCAAGCAGGCGCGCTTTCGCCAGCAAAACCGAAACAGAGCCCGTGTCTG 260
    |||||
Db 247 GCAGCGATCAAGCAGGCGCGCTTTCGCCAGCAAAACCGAAACAGGTTGAAGTTCGCCCA 188
    |||||
```

```
QY 261 GTACTACGTGAAGAGGTCCTCAACAGCAGGCTGCAGCGCTTCTACTCCCTGCGCCA 320
    |||||
Db 187 GGCCACTGTGGCCATGATGAACAGAGGATGAGCTGGAGGAGAGACAGCCCTGGTGA 128
    |||||
```

```
QY 321 C 321
    |
Db 127 C 127
    |
```

## RESULT 10

```
US-10-029-386-8336
```

```
; Sequence 8336, Application US/10029386
; Publication No. US20030194704A1
```

```
; GENERAL INFORMATION:
; APPLICANT: Penn, Sharron G.
```

```
; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
; THE ANALYSIS OF EXPRESSION ANALYSIS TWO
```

```
; FILE REFERENCE: AEOMICA-X-2
; CURRENT APPLICATION NUMBER: US/10/029,386
; CURRENT FILING DATE: 2001-12-20
```

```
; NUMBER OF SEQ ID NOS: 34288
; SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
; SEQ ID NO 8336
; LENGTH: 500
```

```
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: MAP TO CHR16.3
```

```
; OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 1.4
; OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 1.4
; OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 1.2
```



OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 1.5  
OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 1.3  
OTHER INFORMATION: SWISSPROT HIT: Q92883, EVALUE 1.20e+00  
OTHER INFORMATION: NT HIT: g115315787, EVALUE 2.00e-65  
OTHER INFORMATION: EST\_HUMAN HIT: AL135642.1, EVALUE 2.00e-65  
US-10-029-386-8336

Query Match 7.2%; Score 128; DB 17; Length 500;  
Best Local Similarity 100.0%; Pred. No. 1.5e-28;  
Matches 128; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 120 CAGGGTCACCTGCTGTGTGCCCGCTTGAAGCCGCTCTGCAGCATGGCTTGAAGAGGAG 179  
Db |||||||  
Qy 341 CAGGGTCACCTGCTGTGTGCCCGCTTGAAGCCGCTCTGCAGCATGGCTTGAAGAGGAG 400  
Db |||||||  
Qy 180 TCGAGGATTGGCACTCACAGCGGCGGCGATCAAGCAGGCGCGGCTTGGCAGCAAAAC 239  
Db |||||||  
Qy 401 TCGAGGATTGGCACTCACAGCGGCGGCGATCAAGCAGGCGCGGCTTGGCAGCAAAAC 460  
Db |||||||  
Qy 240 CGAAACAG 247  
Db |||||||  
Qy 461 CGAAACAG 468  
Db |||||||

## RESULT 11

US-10-029-386-22036  
Sequence 22036, Application US/10029386  
Publication No. US20030194704A1

GENERAL INFORMATION:  
APPLICANT: Penn, Sharron G.  
APPLICANT: Rank, David R.  
APPLICANT: Hanzel, David K.  
TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR C  
FILE REFERENCE: ASOMICA-X-2  
CURRENT APPLICATION NUMBER: US/10/029,386  
CURRENT FILING DATE: 2001-12-20  
NUMBER OF SEQ ID NOS: 34288  
SOFTWARE: Annonax Sequence Listing Engine vers. 1.1  
SEQ ID NO 22036  
LENGTH: 124  
TYPE: DNA

ORGANISM: Homo sapiens

FEATURE:  
OTHER INFORMATION: MAP TO CHR16.3  
OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 1.4  
OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 1.4  
OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 1.2  
OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 1.5  
OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 1.3  
OTHER INFORMATION: NT HIT: g115315787, EVALUE 9.00e-64  
OTHER INFORMATION: EST\_HUMAN HIT: AL135642.1, EVALUE 1.00e-63  
OTHER INFORMATION: SWISSPROT HIT: Q9SXU1, EVALUE 6.00e+00  
US-10-029-386-22036

Query Match 7.0%; Score 124; DB 17; Length 124;  
Best Local Similarity 100.0%; Pred. No. 1.3e-27;  
Matches 124; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 124 GTACCTGCTGTGTGCCCGCTTGAAGCCGCTCTGCAGCATGGCTTGAAGAGGAGTCGA 183  
Db |||||||  
Qy 1 1 GTACCTGCTGTGTGCCCGCTTGAAGCCGCTCTGCAGCATGGCTTGAAGAGGAGTCGA 60  
Db |||||||  
Qy 184 GGATTGGCACTCACAGCGGCGGCGATCAAGCAGGCGGCGCTTGGCAGCAAAACCGAA 243  
Db |||||||  
Qy 61 GGATTGGCACTCACAGCGGCGGCGATCAAGCAGGCGGCGCTTGGCAGCAAAACCGAA 120  
Db |||||||  
Qy 244 ACAG 247  
Db |||||||  
Qy 121 ACAG 124  
Db |||||||

## RESULT 12

Query Match 4.2%; Score 75; DB 13; Length 633;  
Best Local Similarity 100.0%; Pred. No. 5.2e-12;  
Matches 75; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

US-10-450-763-215/c  
Sequence 215, Application US/10450763  
Publication No. US20050196754A1  
GENERAL INFORMATION:  
APPLICANT: Hyseq, Inc  
TITLE OF INVENTION: NOVEL NUCLEIC ACIDS AND POLYPEPTIDES  
FILE REFERENCE: 790CIP3/US  
CURRENT APPLICATION NUMBER: US/10/450,763  
CURRENT FILING DATE: 2003-06-11  
PRIOR APPLICATION NUMBER: PCT/US01/08631  
PRIOR FILING DATE: 2001-03-30  
PRIOR APPLICATION NUMBER: 09/540,217  
PRIOR FILING DATE: 2000-03-31  
PRIOR APPLICATION NUMBER: 09/649,167  
PRIOR FILING DATE: 2000-08-23  
NUMBER OF SEQ ID NOS: 60736  
SOFTWARE: Custom  
SEQ ID NO 215  
LENGTH: 911  
TYPE: DNA

ORGANISM: Homo sapiens

FEATURE:

NAME/KEY: SIMILAR

LOCATION: (173)..(57)

OTHER INFORMATION: 100% homologous to Homo sapiens Partial sequence of the clone

OTHER INFORMATION: 9 protein, accession number W42400.Smith-Waterman Score=193.

US-10-450-763-215

Query Match 6.7%; Score 119; DB 24; Length 911;  
Best Local Similarity 100.0%; Pred. No. 1.3e-25;  
Matches 119; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 129 CTGTCTGTGTGCCCGCTTGAAGCCGCTCTGCAGCATGGCTTGAAGAGGAGTCGAGGATT 188  
Db |||||||

Qy 174 CTGTCTGTGTGCCCGCTTGAAGCCGCTCTGCAGCATGGCTTGAAGAGGAGTCGAGGATT 115  
Db |||||||

Qy 189 GGCACCTCACAGCGGCGGCGATCAAGCAGGCGGCGCTTGGCAGCAAAACCGAAACAG 247  
Db |||||||

Qy 114 GGCACCTCACAGCGGCGGCGATCAAGCAGGCGGCGCTTGGCAGCAAAACCGAAACAG 56  
Db |||||||

## RESULT 13

US-09-925-065A-681221  
Sequence 681221, Application US/09925065A  
Publication No. US20050238172A9

GENERAL INFORMATION:  
APPLICANT: Wang, David G.  
TITLE OF INVENTION: Identification and Mapping of Single  
FILE REFERENCE: 108827.135  
CURRENT APPLICATION NUMBER: US/09/925,065A  
CURRENT FILING DATE: 2001-08-08  
PRIOR APPLICATION NUMBER: US 60/243,096  
PRIOR FILING DATE: 2000-10-24  
PRIOR APPLICATION NUMBER: US 60/252,147  
PRIOR FILING DATE: 2000-11-20  
PRIOR APPLICATION NUMBER: US 60/250,092  
PRIOR FILING DATE: 2000-11-30  
PRIOR APPLICATION NUMBER: US 60/261,766  
PRIOR FILING DATE: 2001-01-16  
PRIOR APPLICATION NUMBER: US 60/289,846  
PRIOR FILING DATE: 2001-05-09  
NUMBER OF SEQ ID NOS: 957086  
SOFTWARE: Fast-SEQ for Windows Version 4.0  
SEQ ID NO 681221  
LENGTH: 633  
TYPE: DNA

ORGANISM: Homo sapiens

US-09-925-065A-681221

Query Match 4.2%; Score 75; DB 13; Length 633;  
Best Local Similarity 100.0%; Pred. No. 5.2e-12;  
Matches 75; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 427 AGCACTTTTATGAGACTGGTCTTTTGTGATGGATGAAGAAAGGTCCAGTATGCTTCT 486  
Db |||||||  
484 AGCACTTTTATGAGACTGGTCTTTTGTGATGGATGAAGAAAGGTCCAGTATGCTTCT 543  
QY |||||||  
487 ACCATGGCAGCAGGT 501  
Db |||||||  
544 ACCATGGCAGCAGGT 558

RESULT 14  
US-11-097-143-11288  
; Sequence 11288, Application US/11097143  
; Publication No. US2005020858A1  
; GENERAL INFORMATION:  
; APPLICANT: Venter, J. Craig  
; APPLICANT: et al.  
; TITLE OF INVENTION: DETECTION KIT, SUCH AS NUCLEIC ACID  
; TITLE OF INVENTION: ARRAYS, FOR DETECTING EXPRESSION OF 10,000 OR MORE  
; TITLE OF INVENTION: DROSOPHILA GENES.  
; FILE REFERENCE: CL000728  
; CURRENT APPLICATION NUMBER: US/11/097,143  
; CURRENT FILING DATE: 2005-04-04  
; PRIOR APPLICATION NUMBER: 60/157,832  
; PRIOR FILING DATE: 1999-10-05  
; PRIOR APPLICATION NUMBER: 60/160,191  
; PRIOR FILING DATE: 1999-10-19  
; PRIOR APPLICATION NUMBER: 60/161,932  
; PRIOR FILING DATE: 1999-10-28  
; PRIOR APPLICATION NUMBER: 60/164,769  
; PRIOR FILING DATE: 1999-11-12  
; PRIOR APPLICATION NUMBER: 60/173,383  
; PRIOR FILING DATE: 1999-12-28  
; PRIOR APPLICATION NUMBER: 60/175,693  
; PRIOR FILING DATE: 2000-01-12  
; PRIOR APPLICATION NUMBER: 60/184,831  
; PRIOR FILING DATE: 2000-02-24  
; PRIOR APPLICATION NUMBER: 60/191,637  
; PRIOR FILING DATE: 2000-03-23  
; NUMBER OF SEQ ID NOS: 43008  
; SOFTWARE: FastSeq for Window Version 4.0  
; SEQ ID NO 11288  
; LENGTH: 1796  
; TYPE: DNA  
; ORGANISM: DROSOPHILA

US-11-097-143-11288  
Query Match 3.7%; Score 66.8; DB 26; Length 1796;  
Best Local Similarity 49.4%; Pred. No. 3.1e-09;  
Matches 173; Conservative 0; Mismatches 177; Indels 0; Gaps 0;  
QY 233 GCAAAACCGAAACAGAGCCGCTGTTCTGTACTACGTGAAGAGGTCTTCAACAAGCAGC 292  
Db |||||||  
396 GCAGCAACAACATGACGCCACTTTCTGGGAATCTGCCAGACGCATTTGACGCCCCACG 455  
QY 293 AGCTGACGCCCTTCTACTCCCTGGGCCACATCGCTCAGACGTGGCGGGGTGCGCCT 352  
Db |||||||  
456 AACCTCAGAGGTATATGGACCTTTAAGCAGATCTGGACGAATGTGGCAGAGCAGTGCCT 515  
QY 353 GGCTGGCTGTGCTTCAAGAACACTCCCTGGAGCGCTACCTGCACATGCTCTCGGCCG 412  
Db |||||||  
516 TCATACCGGCCCACTGAACAGAGCACTGATAGCCATGTCTCACCTGGCTGAGCG 575  
QY 413 ACCGCTGACGCTGAGCAGCTTTTATGAAGACTGGTCTTTGTGATGGATGAAGAAAGGT 472  
Db |||||||  
576 ACAGGAGCAGCTGCATCGATTTTACACACTTGGTCCCTGCTACTCAACGATGAGCGG 635  
QY 473 CCAGTATGCTTCTTACCATCGCAGAGTCTGAACTCCATCTCTTTGCGATTAAACATCG 532  
Db |||||||  
636 CCAGAGAGCTGCCAGAGATCGTAGACTCCCTCAGTGTACTGTTCGCCCTCAATGTGG 695  
QY 533 ACAACAGAGATTTGAACGGGCGAGAGTAAGTTTGTCTCCACCGTTTCAGAC 582

Search completed: October 26, 2005, 01:43:40  
Job time : 1484 secs

Db 696 ACACCACTGAGCTTAATGCACCCAGGAGATCGACCCCAAGTGTTCGCCGTC 745

## RESULT 15

US-11-097-143-11287/c  
; Sequence 11287, Application US/11097143  
; Publication No. US2005020858A1  
; GENERAL INFORMATION:  
; APPLICANT: Venter, J. Craig  
; APPLICANT: et al.  
; TITLE OF INVENTION: DETECTION KIT, SUCH AS NUCLEIC ACID  
; TITLE OF INVENTION: ARRAYS, FOR DETECTING EXPRESSION OF 10,000 OR MORE  
; TITLE OF INVENTION: DROSOPHILA GENES.  
; FILE REFERENCE: CL000728  
; CURRENT APPLICATION NUMBER: US/11/097,143  
; CURRENT FILING DATE: 2005-04-04  
; PRIOR APPLICATION NUMBER: 60/157,832  
; PRIOR FILING DATE: 1999-10-05  
; PRIOR APPLICATION NUMBER: 60/160,191  
; PRIOR FILING DATE: 1999-10-19  
; PRIOR APPLICATION NUMBER: 60/161,932  
; PRIOR FILING DATE: 1999-10-28  
; PRIOR APPLICATION NUMBER: 60/164,769  
; PRIOR FILING DATE: 1999-11-12  
; PRIOR APPLICATION NUMBER: 60/173,383  
; PRIOR FILING DATE: 1999-12-28  
; PRIOR APPLICATION NUMBER: 60/175,693  
; PRIOR FILING DATE: 2000-01-12  
; PRIOR APPLICATION NUMBER: 60/184,831  
; PRIOR FILING DATE: 2000-02-24  
; PRIOR APPLICATION NUMBER: 60/191,637  
; PRIOR FILING DATE: 2000-03-23  
; NUMBER OF SEQ ID NOS: 43008  
; SOFTWARE: FastSeq for Windows Version 4.0  
; SEQ ID NO 11287  
; LENGTH: 3796  
; TYPE: DNA  
; ORGANISM: DROSOPHILA

US-11-097-143-11287  
Query Match 3.7%; Score 66.8; DB 26; Length 3796;  
Best Local Similarity 49.4%; Pred. No. 4.5e-09;  
Matches 173; Conservative 0; Mismatches 177; Indels 0; Gaps 0;  
QY 233 GCAAAACCGAAACAGAGCCGCTGTTCTGTACTACGTGAAGAGGTCTTCAACAAGCAGC 292  
Db |||||||  
2401 GCAGCAACAACATGACGCCACTTTCTGGGAATTTGCCAGACGCATTTGACGCCCCACG 2342  
QY 293 AGCTGACGCCCTTCTACTCCCTGGGCCACATCGCTCAGACGTGGCGGGGTGCGCCT 352  
Db |||||||  
2341 AACGTGAGAGGTATATGGACCTTTAAGCAGATCTGGACGAATGTGGCAGAGACGTGCCT 2282  
QY 353 GGCTGGCTGTGCTTCAAGAACACTCCCTGGAGCGCTACCTGCACATGCTCTCTGGCCG 412  
Db |||||||  
2281 TCATACCGGCCCACTGAACAGAGCACTGATAGCCATGTCTCACCTGGCTGAGCG 2222  
QY 413 ACCGCTGACGCTGAGCAGCTTTTATGAAGACTGGTCTTTGTGATGGATGAAGAAAGGT 472  
Db |||||||  
2221 ACAGGAGCAGCTGCATCGATTTTACACACTTGGTCCCTGCTACTCAACGATGAGCGG 2162  
QY 473 CCAGTATGCTTCTTACCATGGCAGAGTCTGAACTCCATCTCTTTGCGATTAAACATCG 532  
Db |||||||  
2161 CCAGAGAGCTGCCAGAGATCGTAGACTCCCTCAGTGTACTGTTCGCCCTCAATGTGG 2102  
QY 533 ACAACAGAGATTTGAACGGGCGAGAGTAAGTTTGTCTCCACCGTTTCAGAC 582  
Db |||||||  
2101 ACACCACTGAGCTTAATGCACCCAGGAGATCGACCCCAAGTGTTCGCCGTC 2052

GenCore version 5.1.6  
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OM nucleic - nucleic search, using sw model

Run on: October 25, 2005, 18:28:14 ; Search time 286 Seconds  
(without alignments)  
10195.263 Million cell updates/sec

Title: US-10-070-255-4  
Perfect score: 1782  
Sequence: 1 atgagcgatcacagacaa.....taaatgtcttcattccttg 1782

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 1202784 seqs, 818138359 residues

Total number of hits satisfying chosen parameters: 2405568

Minimum DB seq length: 0  
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : Issued Patents, NA.\*  
1: /cgn2\_6/ptodata/1/ina/5A\_COMB.seq.\*  
2: /cgn2\_6/ptodata/1/ina/5B\_COMB.seq.\*  
3: /cgn2\_6/ptodata/1/ina/6A\_COMB.seq.\*  
4: /cgn2\_6/ptodata/1/ina/6B\_COMB.seq.\*  
5: /cgn2\_6/ptodata/1/ina/PTUS\_COMB.seq.\*  
6: /cgn2\_6/ptodata/1/ina/backfiles.seq.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	133.4	7.5	190	4	US-09-513-999C-16002
2	66.8	3.7	811	4	US-09-270-767-12402
3	56.8	3.2	7218	1	US-08-232-463-14
4	54.4	3.1	36731	4	US-09-949-016-13770
5	54	3.0	601	4	US-09-949-016-13770
6	52.8	3.0	476	4	US-09-573-080A-118
7	52.2	2.9	961	4	US-09-976-594-221
8	49.6	2.8	9811	4	US-09-949-016-12759
9	47	2.6	100	4	US-09-513-999C-16212
10	46.8	2.6	60276	4	US-09-949-016-15004
11	46.8	2.6	60338	4	US-09-949-016-15094
12	46.8	2.6	94133	4	US-09-949-016-11901
13	46.8	2.6	94133	4	US-09-949-016-12713
14	46.8	2.6	94135	4	US-09-949-016-15934
15	46.8	2.6	94135	4	US-09-949-016-15935
16	46.8	2.6	94135	4	US-09-949-016-15936
17	46.8	2.6	94135	4	US-09-949-016-15937
18	44.8	2.5	474	4	US-09-621-976-18033
19	43.2	2.4	1002	4	US-09-902-540-8383
20	43.2	2.4	7035	4	US-09-902-540-878
21	43.2	2.4	23802	4	US-09-949-016-12107
22	43.2	2.4	23803	4	US-09-949-016-15878
23	43	2.4	1304	4	US-09-902-540-2596
24	43	2.4	16584	4	US-09-902-540-1119
25	42	2.4	23417	4	US-09-902-540-1207
26	41.4	2.3	505	4	US-09-621-976-15639
27	41.2	2.3	1926	3	US-09-249-585A-2

28	41.2	2.3	1926	4	US-09-410-399-3	Sequence 3, Appli
29	41.2	2.3	2580	3	US-09-050-863-2	Sequence 2, Appli
30	41.2	2.3	2580	3	US-09-359-081-2	Sequence 2, Appli
31	41.2	2.3	5452	2	US-09-130-114-1	Sequence 1, Appli
32	41.2	2.3	8705	4	US-09-647-344A-14	Sequence 14, Appli
33	41.2	2.3	9600	3	US-08-910-647-1	Sequence 1, Appli
34	41.2	2.3	9600	3	US-09-620-925-1	Sequence 1, Appli
35	41.2	2.3	10596	1	US-07-884-811-15	Sequence 15, Appli
36	41.2	2.3	10596	1	US-07-885-971-15	Sequence 15, Appli
37	41.2	2.3	10596	1	US-08-087-783A-15	Sequence 15, Appli
38	41.2	2.3	10596	1	US-08-194-088B-15	Sequence 15, Appli
39	41.2	2.3	10596	2	US-08-194-087-15	Sequence 15, Appli
40	41.2	2.3	10596	5	PCT-US93-04648-15	Sequence 15, Appli
41	41.2	2.3	16080	4	US-09-724-566A-48	Sequence 48, Appli
42	41.2	2.3	16080	4	US-09-471-669A-48	Sequence 48, Appli
43	41	2.3	1275	4	US-09-252-931A-251	Sequence 251, App
44	41	2.3	1785	4	US-09-252-931A-251	Sequence 281, App
45	40.6	2.3	1132	3	US-09-370-838-256	Sequence 256, App

ALIGNMENTS

RESULT 1

US-09-513-999C-16002  
; Sequence 16002, Application US/09513999C  
; Patent No. 6783961  
; GENERAL INFORMATION:  
; APPLICANT: Dumas Milne Edwards, J.B.  
; APPLICANT: Duclert, A.  
; APPLICANT: Giordano, J.Y.  
; TITLE OF INVENTION: Expressed Sequence Tags and Encoded Human Proteins.  
; PATENT NO. 6783961  
; FILE REFERENCE: 59, US2, REG  
; CURRENT APPLICATION NUMBER: US/09/513,999C  
; CURRENT FILING DATE: 2000-02-24  
; PRIOR APPLICATION NUMBER: US 60/122,487  
; PRIOR FILING DATE: 1999-02-26  
; NUMBER OF SEQ ID NOS: 36681  
; SOFTWARE: Patent.pm  
; SEQ ID NO 16002  
; LENGTH: 190  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-09-513-999C-16002

Query Match	7.5%	Score 133.4;	DB 4;	Length 190;
Best Local Similarity	99.3%	Pred. No. 8.4e-26;		
Matches 134;	Conservative 0;	Mismatches 1;	Indels 0;	Gaps 0;
Qy	1464	CAGATCACTGCGAAACCTGCTCGACGGTGAGATGGAGCAGCTCAGCGGGCTCCGGCAAGA	1523	
Db	19	CAGATCACTGCGAAACCTGCTCGACGGTGAGATGGAGCAGCTCAGCGGGCTCCGGCAAGA	78	
Qy	1524	GGTGGACACCTTGAAGAGGAGGTGGCTGAACAGGAGCGGAGGCGCATGAAGGTCCA	1583	
Db	79	GGTGGACACCTTGAAGAGGAGGTGGCTGAACAGGAGCGGAGGCGCATGAAGGTCCA	138	
Qy	1584	GGCGCTGCCAGCTA	1598	
Db	139	GGCGCTGCCAGGTA	153	

RESULT 2

US-09-270-767-12402  
; Sequence 12402, Application US/09270767  
; Patent No. 6703491  
; GENERAL INFORMATION:  
; APPLICANT: Homburger et al.  
; TITLE OF INVENTION: Nucleic acids and proteins of Drosophila melanogaster  
; FILE REFERENCE: File Reference: 7326-094  
; CURRENT APPLICATION NUMBER: US/09/270,767  
; CURRENT FILING DATE: 1999-03-17



```

RESULT 6
US-09-573-080A-118
; Sequence 118, Application US/09573080A
; Patent No. 6828097
; GENERAL INFORMATION:
; APPLICANT: JOAN, KNOLL
; APPLICANT: ROCAN, PETER
; TITLE OF INVENTION: SINGLE COPY GENOMIC HYBRIDIZATION PROBES AND METHOD OF GENERATING
; FILE REFERENCE: 30307
; CURRENT APPLICATION NUMBER: US/09/573,080A
; CURRENT FILING DATE: 2000-05-16
; NUMBER OF SEQ ID NOS: 479
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 118
; LENGTH: 476
; TYPE: DNA

```

```

RESULT 7
US-09-976-594-221
; Sequence 221, Application US/09976594
; Patent No. 6673549
; GENERAL INFORMATION:
; APPLICANT: Furness, Michael
; APPLICANT: Buchbinder, Jenny
; TITLE OF INVENTION: GENES EXPRESSED
; FILE REFERENCE: PA-0041 US
; CURRENT APPLICATION NUMBER: US/09/976
; CURRENT FILING DATE: 2001-10-12
; PRIOR APPLICATION NUMBER: 60/240,409
; PRIOR FILING DATE: 2000-10-12
; NUMBER OF SEQ ID NOS: 1143
; SOFTWARE: PERL Program
; SEQ ID NO 221
; LENGTH: 961
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; OTHER INFORMATION: Incyte ID No. 66
US-09-976-594-221

```

	Query Match	2.9%	Score 52.2;	DB 4;	Length 961;
	Best Local Similarity	71.1%;	Pred. No. 0.0009;		
	Matches	69;	Conservative	0;	Mismatches 28; Indels 0; Gaps 0;
Qy	1679	GTTCCTGATAGTCTCATTTGAGCTCTCGAGTCCAGTCCTTTCCCTGAAGCTGTGTTCTCT	1738		
Db	861	GTCCCTGGTGACATCTTTTGAGCCACTAGACCAAGCTTTACCTGAAGCAGAGCTACCTCA	920		
Qy	1739	GGACTTTTCATGTATGTGAGCCCAATAATGCTTTCA	1775		
Db	921	GAGCTTTTCAGCTATGTGAGCCCAATAAATCTGTCA	957		

## RESULT 8

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US-09-949-016-12759
; Sequence 12759, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; PRIOR FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 12759
; LENGTH: 9811
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-12759

Query Match 2.8%; Score 49.6; DB 4; Length 9811;
Best Local Similarity 61.7%; Pred. No. 0.013; Mismatches 49; Indels 0; Gaps 0;
Matches 79; Conservative 0;

QY 1654 AATGGAGGAGAGAGAAATCCACAGTTCCTGATAGTCTCATTTGAGCTCCTGGATCCAG 1713
DB 3752 AAGAAGAGAGTGTGAGTGACCATTTCTTATTCATCCTTTGAGCCCTACACCTAA 3811

QY 1714 TCCTTCTGAAGCTGTGTTTCTCTGGACTTTTCATGTATGTGAGCCAAATAATGCTTT 1773
DB 3812 TCATTCATGAAGCTATACCACTCCAGCCCTTTCAATTACTTGAGCCAAATTCCTT 3871

QY 1774 CATTCCTT 1781
DB 3872 TAATAGTT 3879

RESULT 9
US-09-513-999C-16212
; Sequence 16212, Application US/09513999C
; Patent No. 6783961
; GENERAL INFORMATION:
; APPLICANT: Dumas Milne Edwards, J.B.
; APPLICANT: Duclert, A.
; APPLICANT: Giordano, J.Y.
; TITLE OF INVENTION: Expressed Sequence Tags and Encoded Human Proteins.
; Patent No. 6783961
; FILE REFERENCE: 59.US2.REG
; CURRENT APPLICATION NUMBER: US/09/513,999C
; CURRENT FILING DATE: 2000-02-24
; PRIOR APPLICATION NUMBER: US 60/122,487
; PRIOR FILING DATE: 1999-02-26
; NUMBER OF SEQ ID NOS: 36681
; SOFTWARE: Patent.pm
; SEQ ID NO 16212
; LENGTH: 100
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 13
; OTHER INFORMATION: m=a or c
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: 14
; OTHER INFORMATION: m=a or c
US-09-513-999C-16212

Query Match 2.6%; Score 47; DB 4; Length 100;

US-09-949-016-15004/c
; Sequence 15004, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 15004
; LENGTH: 60276
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(60276)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-15004

Query Match 2.6%; Score 46.8; DB 4; Length 60276;
Best Local Similarity 73.2%; Pred. No. 0.17; Mismatches 22; Indels 0; Gaps 0;
Matches 60; Conservative 0;

QY 1692 TCATTGAGCTCCTGGATCCAGTCTTTCTCTGAAGCTGTGTTTCTCTGGACTTTTCATGT 1751
DB 10119 TCTTAGAGCACCTGATCCAGCCCTGCTCTGAAGCCATCTACCCCTGGACTTCTCAGTT 10060

QY 1752 ATGTGAGCCAAATAATGCTTT 1773
DB 10059 GTGTGAGCCAAGAAATGCTTT 10038

RESULT 11
US-09-949-016-15694/c
; Sequence 15694, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 15694
; LENGTH: 60338
; TYPE: DNA
; ORGANISM: Human
```

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; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (1)...(60338)
; OTHER INFORMATION: n = A,T,C or G
US-09-949-016-15694

Query Match          2.6%; Score 46.8; DB 4; Length 60338;
Best Local Similarity 73.2%; Pred. No. 0.17;
Matches 60; Conservative 0; Mismatches 22; Indels 0; Gaps 0;

Qy 1692 TCATTGAGCTCCGATCAGTCTTTCTCGAAGCTGTGTTCTCTCGACTTTTCAGT 1751
Db 10181 TCTTAGAAGCACCTGGATCCAGCCCTGCTGAAGCCATCTACCCCTGGACTTTCAGT 10122

Qy 1752 ATGTGAGCCAATAATTCCTTT 1773
Db 10121 GTGTGAGCCAAGATTGTCCTT 10100

RESULT 12
US-09-949-016-11901/c
; Sequence 11901, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 11901
; LENGTH: 94133
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-11901

Query Match          2.6%; Score 46.8; DB 4; Length 94133;
Best Local Similarity 67.3%; Pred. No. 0.21;
Matches 66; Conservative 0; Mismatches 32; Indels 0; Gaps 0;

Qy 1678 AGTTCCTGATAGTCTCATTTGAGCTCCTGGATCCAGTCTTTCTCGAAGCTGTGTTCTC 1737
Db 40227 AGAGCTGAGAACACAGTTTAAGCATTTGGAGTCAGCTATGCTCGAAGCCATGCTCCCTC 40168

Qy 1738 TGGACTTTTCATGTATGTGAGCCCAATAAATGCTTTCA 1775
Db 40167 TAGACTTTTCAGTTACATGAGCCCAATAAATATTTT 40130

RESULT 13
US-09-949-016-12713/c
; Sequence 12713, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
```

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; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 12713
; LENGTH: 94133
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-12713

Query Match          2.6%; Score 46.8; DB 4; Length 94133;
Best Local Similarity 67.3%; Pred. No. 0.21;
Matches 66; Conservative 0; Mismatches 32; Indels 0; Gaps 0;

Qy 1678 AGTTCCTGATAGTCTCATTTGAGCTCCTGGATCCAGTCTTTCTCGAAGCTGTGTTCTC 1737
Db 40227 AGAGCTGAGAACACAGTTTAAGCATTTGGAGTCAGCTATGCTCGAAGCCATGCTCCCTC 40168

Qy 1738 TGGACTTTTCATGTATGTGAGCCCAATAAATGCTTTCA 1775
Db 40167 TAGACTTTTCAGTTACATGAGCCCAATAAATATTTT 40130

RESULT 14
US-09-949-016-15934/c
; Sequence 15934, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 15934
; LENGTH: 94135
; TYPE: DNA
; ORGANISM: Human
US-09-949-016-15934

Query Match          2.6%; Score 46.8; DB 4; Length 94135;
Best Local Similarity 67.3%; Pred. No. 0.21;
Matches 66; Conservative 0; Mismatches 32; Indels 0; Gaps 0;

Qy 1678 AGTTCCTGATAGTCTCATTTGAGCTCCTGGATCCAGTCTTTCTCGAAGCTGTGTTCTC 1737
Db 40227 AGAGCTGAGAACACAGTTTAAGCATTTGGAGTCAGCTATGCTCGAAGCCATGCTCCCTC 40168

Qy 1738 TGGACTTTTCATGTATGTGAGCCCAATAAATGCTTTCA 1775
Db 40167 TAGACTTTTCAGTTACATGAGCCCAATAAATATTTT 40130

RESULT 15
US-09-949-016-15935/c
; Sequence 15935, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
; TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
; CURRENT APPLICATION NUMBER: US/09/949,016
; CURRENT FILING DATE: 2000-04-14
; PRIOR APPLICATION NUMBER: 60/241,755
; PRIOR FILING DATE: 2000-10-20
```

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; PRIOR APPLICATION NUMBER: 60/237,768
; PRIOR FILING DATE: 2000-10-03
; PRIOR APPLICATION NUMBER: 60/231,498
; PRIOR FILING DATE: 2000-09-08
; NUMBER OF SEQ ID NOS: 207012
; SOFTWARE: FASTSEQ for Windows Version 4.0
; SEQ ID NO 15935
; LENGTH: 94135
; TYPE: DNA
; ORGANISM: Human
; US-09-949-016-15935

Query Match      2.6%; Score 46.8; DB 4; Length 94135;
Best Local Similarity 67.3%; Pred. No. 0.21;
Matches 66; Conservative 0; Mismatches 32; Indels 0; Gaps 0;

Qy      1678 AGTTCCTGATAGTCTCATTTTGAGCTCCCTGGATCCAGTCTTTTCCTGAAGCTGTGTTTCTCT 1737
          |||||
Db      40227 AGAGCCTGAGACACACAGTTTTAAGCATTTTGAGTTCAGTATGCTTGAGCCCATGCTCCCTC 40168
          |||||

Qy      1738 TGGACITTTTCATGTATGTGAGGCAATAATTCCTTTCA 1775
          |||||
Db      40167 TAGACITTTTCAGTTACATGAGGCAATAAATAATTTTA 40130
          |||||

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Search completed: October 25, 2005, 21:20:34  
Job time : 289 secs



GenCore version 5.1.6  
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OM nucleic - nucleic search, using sw model

Run on: October 25, 2005, 21:16:28 ; Search time 981 Seconds  
(without alignments)  
10753.292 Million cell updates/sec

Title: US-10-070-255-4  
Perfect score: 1782  
Sequence: 1 atgagcggatcacagacaa.....taaatgcttcttcattccttg 1782

Scoring table: IDENTITY NUC  
Gapop 10.0 , Gapext 1.0

Searched: 4390206 seqs, 2959870667 residues

Total number of hits satisfying chosen parameters: 8780412

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : N\_Geneseq\_16Dec04:\*

- 1: Geneseqn1980s:\*
- 2: Geneseqn1990s:\*
- 3: Geneseqn2000s:\*
- 4: Geneseqn2001as:\*
- 5: Geneseqn2001bs:\*
- 6: Geneseqn2002as:\*
- 7: Geneseqn2002bs:\*
- 8: Geneseqn2003as:\*
- 9: Geneseqn2003bs:\*
- 10: Geneseqn2003cs:\*
- 11: Geneseqn2003ds:\*
- 12: Geneseqn2004as:\*
- 13: Geneseqn2004bs:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	1782	100.0	1782	4	AAD04333
2	1742	97.8	1906	2	AAV03323
3	1595	89.5	2873	4	AAD04335
4	1595	89.5	3139	4	AAD04334
5	1129.8	63.4	2248	4	AAH15665
6	816.8	45.8	6045	4	AAH18235
7	639.8	35.9	7330	4	AAH04973
8	364.2	20.4	33147	4	AAK67282
9	275.8	15.5	280	6	ABS72233
10	247	13.9	454	5	AAS64410
11	227.8	12.8	542	12	ACH75347
12	224.8	12.6	284	12	ACH89047
13	207.6	11.6	463	5	AAS92200
14	135.4	7.6	474	5	AAS73549
15	133.4	7.5	190	3	AAC11927
16	128	7.2	500	12	ACH75141
17	124	7.0	124	12	ACH88841
18	119	6.7	911	5	AAS64411
19	66.8	3.7	1796	4	ABL09365
20	66.8	3.7	3796	4	ABL09364

C 21	63.8	3.6	3931	4	ABA09172	ABA09172 Human BG3
C 22	59.6	3.3	214520	10	ADL13471	Adl13471 Osteoarthritis
C 23	59	3.3	3927	10	ADP81729	Adf81729 Leukaemia
C 24	58.4	3.3	595	13	ADOS3383	Adqs3383 Novel can
C 25	54.4	3.1	813	5	AAS55362	Aas55362 DNA encod
C 26	54	3.0	464	9	ACH27965	Ach27965 Human adu
C 27	52.8	3.0	263	6	ABL86718	Ab186718 Human ova
C 28	52.8	3.0	476	7	ADS31085	Ads31085 Human gen
C 29	52.2	2.9	729	10	ADE07225	Ade07225 Novel Cod
C 30	52.2	2.9	941	5	ABA21208	Aba21208 Human ner
C 31	52.2	2.9	941	5	ABA21210	Aba21210 Human ner
C 32	52.2	2.9	961	12	ADL12492	Adl12492 Human ste
C 33	52.2	2.9	3916	13	ADP07577	Adp07577 Full leng
C 34	51.8	2.9	617	5	ABA14117	Aba14117 Human ner
C 35	50.2	2.8	2000	8	ADA71938	Ada71938 Rice gene
C 36	50	2.8	371	9	ACH37072	Ach37072 Human end
C 37	50	2.8	446	6	ABN95020	Abn95020 Gene #151
C 38	50	2.8	769	4	AAI95747	Aai95747 Human neu
C 39	50	2.8	5684	3	AAC59953	Aac59953 Human sec
C 40	50	2.8	5684	8	ABZ73290	Abz73290 Secreted
C 41	50	2.8	5684	10	ABZ66903	Abz66903 Human sec
C 42	50	2.8	7093	6	ABZ35416	Abz35416 Human gen
C 43	50	2.8	10126	4	AAI99260	Aai99260 Human exc
C 44	50	2.8	10126	5	AAI63610	Aai63610 Human kid
C 45	49.6	2.8	90597	10	ADJ72363	Adj72363 Streptomy

ALIGNMENTS

RESULT 1

AAD04333

ID AAD04333 standard; cDNA; 1782 BP.

XX AAD04333;

DT 04-JUL-2001 (first entry)

DE Human TNF receptor-associated factor (TRAF2) binding protein, IREN cDNA.

XX Human; Tumour Necrosis Factor; TNF; TNF Receptor Associated Factor;

XX TRAF2; TRAF2 binding protein; IREN; IkappaB Regulator; immunosuppressive;

XX nuclear factor-kappaB; NF-kappaB; cytostatic; tumour; AIDS;

XX acquired immune deficiency syndrome; rheumatic disease; apoptosis;

XX autoimmune disease; septic shock; graft-vs-host reaction; inflammation;

XX anorexia; anti-HIV; therapy; ss.

XX Homo sapiens.

OS Key

FT CDS

FT Location/Qualifiers

FT 1..1626

FT /\*tag= a

FT /product= "Human TRAF2-binding protein, IREN"

FT

XX WO200116314-A1.

XX 08-MAR-2001.

XX 31-AUG-2000; 2000WO-IL000517.

XX 02-SEP-1999; 99IL-00131719.

XX (VEDA ) YEDA RES & DEV CO LTD.

XX Wallach D, Malinin N, Sinha I, Leu S;

XX WPI; 2001-281387/29.

XX P-PSDB; AAE00683.

XX New DNA sequence encoding Tumor Necrosis Factor receptor associated

XX factor (TRAF) binding proteins (IREN) for treatment or prevention of

XX pathological conditions associated with NF-kappaB induction.

XX

Claim 1; Fig 3B; 118bp; English.

PS The present cDNA sequence encodes human tumour necrosis factor (TNF)  
 CC receptor-associated factor (TRAF2) binding protein, IREN. A fragment of  
 CC this IREN molecule is capable of binding to TRAF2 protein at position 225  
 CC -501. The invention relates to human tumour necrosis factor (TNF)  
 CC receptor-associated factor (TRAF2) binding protein designated as IREN  
 CC (IkappaB Regulator), its isoforms IREN-10B, IREN-E and their  
 CC corresponding cDNA molecules. IREN is useful for modulating/mediating the  
 CC activity of transcription factor NF (Nuclear Factor)-kappaB or any other  
 CC intracellular signalling activity mediated by TRAF2. IREN is useful in  
 CC the prevention and treatment of a pathological condition associated with  
 CC NF-kappaB induction (abnormal) e.g. AIDS (acquired immune deficiency  
 CC syndrome), autoimmune diseases, tumours, rheumatic diseases, anorexia,  
 CC septic shock and graft-vs-host reactions. IREN also plays an important  
 CC role in the control of inflammation and other non-apoptotic effects of  
 CC TNF as well as in the control of apoptosis. The invention also relates to  
 CC method for screening, identifying and producing a molecule capable of  
 CC modulating activities mediated by IREN. IREN antibodies are useful for  
 CC the purification of new proteins from different sources, including cell  
 CC extracts or transformed cell lines, in addition IREN can be used in  
 CC diagnostic purposes for identifying disorders related to abnormal  
 CC functioning of cellular effects mediated directly by TRAF proteins  
 CC  
 XX  
 SQ Sequence 1782 BP; 456 A; 458 C; 504 G; 364 T; 0 U; 0 Other;

Query Match 100.0%; Score 1782; DB 4; Length 1782;  
 Best Local Similarity 100.0%; Pred. No. 0;  
 Matches 1782; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATGAGCGGATCACAGAACATGACAAAGACAATTTCTGCTGAGCGGACTGCTGGATGCA 60  
 DB 1 ATGAGCGGATCACAGAACATGACAAAGACAATTTCTGCTGAGCGGACTGCTGGATGCA 60

QY 61 GTGAAACAGTGCCAGATCCCGTTTGGAGGAGAAAGAGATTGCTCGGATCCGACAGC 120  
 DB 61 GTGAAACAGTGCCAGATCCCGTTTGGAGGAGAAAGAGATTGCTCGGATCCGACAGC 120

QY 121 AGGGTCACTGTCTGTGTCGCCAGTTTGAAGCCGTCCTGAGCATGGCTTTGAAGAGAGT 180  
 DB 121 AGGGTCACTGTCTGTGTCGCCAGTTTGAAGCCGTCCTGAGCATGGCTTTGAAGAGAGT 180

QY 181 CGAGGATTGGCACTCACAGCGGAGCGATCAAGCAGCGCGGCTTTGCGACGAAACC 240  
 DB 181 CGAGGATTGGCACTCACAGCGGAGCGATCAAGCAGCGCGGCTTTGCGACGAAACC 240

QY 241 GAAACAGAGCCCTGTTTGTGTTACTAGTGAAGAGGTCTTCAACAGCAGAGTGCAG 300  
 DB 241 GAAACAGAGCCCTGTTTGTGTTACTAGTGAAGAGGTCTTCAACAGCAGAGTGCAG 300

QY 301 CGCTTCTACTCCCTGCGCCACATCGCTCAGACGTGGCGCGGGTCCGCGCTGGCTGCGC 360  
 DB 301 CGCTTCTACTCCCTGCGCCACATCGCTCAGACGTGGCGCGGGTCCGCGCTGGCTGCGC 360

QY 361 TGTGCCCTCAACGAAACACTCCCTGGAGCGCTACTGTCACATGCTCCTGGCGGACCGCTGC 420  
 DB 361 TGTGCCCTCAACGAAACACTCCCTGGAGCGCTACTGTCACATGCTCCTGGCGGACCGCTGC 420

QY 421 AGGCTGAGCACTTTTATGAAGACTGGTCTTTTGTGATGATGAAGAAAGGTCCAGTATG 480  
 DB 421 AGGCTGAGCACTTTTATGAAGACTGGTCTTTTGTGATGATGAAGAAAGGTCCAGTATG 480

QY 481 CTTTCTACCATGCGAGAGTCTGAACTCCATCTCTTTGCGATTAACTCGACCAACAG 540  
 DB 481 CTTTCTACCATGCGAGAGTCTGAACTCCATCTCTTTGCGATTAACTCGACCAACAG 540

QY 541 GATTTGAAACGGGAGAGTAAAGTTTGTCTCCACCGTTTTCAGACCTCTTAAAGAGTCAACG 600  
 DB 541 GATTTGAAACGGGAGAGTAAAGTTTGTCTCCACCGTTTTCAGACCTCTTAAAGAGTCAACG 600

QY 601 CAGAACGTGACCTCTTCTGCTGAAGAGTCCACGCAAGAGTGAAGAGCTGTTTCAGGGAG 660  
 DB 601 CAGAACGTGACCTCTTCTGCTGAAGAGTCCACGCAAGAGTGAAGAGCTGTTTCAGGGAG 660

QY 661 ATCAGAGCCTCCTCTGCGCTCTCCATCTCATCAAAACCTGAAACAGAGACCGACCCCTTG 720  
 DB 661 ATCAGAGCCTCCTCTGCGCTCTCCATCTCATCAAAACCTGAAACAGAGACCGACCCCTTG 720

QY 721 CTTGTCGTGTCCAGGAATGTGAGTGTGATGCCAAATGCAAAAGAGCGGAGAGAGAAA 780  
 DB 721 CTTGTCGTGTCCAGGAATGTGAGTGTGATGCCAAATGCAAAAGAGCGGAGAGAGAAA 780

QY 781 AAGAAAGTGACCAACAATAATCTCATTTGATGATGAGAAAGATGAGAGCAAACTCCGACCCCTCT 900  
 DB 781 AAGAAAGTGACCAACAATAATCTCATTTGATGATGAGAAAGATGAGAGCAAACTCTCGGGAC 840

QY 841 GTGTTTAAAAAGACACTCGGGCAGGGAGAGCTCAGAGCAAACTCCGACCCCTCTCT 900  
 DB 841 GTGTTTAAAAAGACACTCGGGCAGGGAGAGCTCAGAGCAAACTCCGACCCCTCTCT 900

QY 901 GTCAATATCATGTCCGCCCTTTGAAAGCCCTTTCGGGCCCTAACTCCAAATGGAAGTCAGAGC 960  
 DB 901 GTCAATATCATGTCCGCCCTTTGAAAGCCCTTTCGGGCCCTAACTCCAAATGGAAGTCAGAGC 960

QY 961 AGCAACTCATGGAATAATTGATTCCTCTTTGAAACGGGGAGTTTGGGTACCAAGAGCTT 1020  
 DB 961 AGCAACTCATGGAATAATTGATTCCTCTTTGAAACGGGGAGTTTGGGTACCAAGAGCTT 1020

QY 1021 GATGTGAAAGCATCGATGATGAGATGTGGATGAAACCGAAGATGACGTGTATGGAAC 1080  
 DB 1021 GATGTGAAAGCATCGATGATGAGATGTGGATGAAACCGAAGATGACGTGTATGGAAC 1080

QY 1081 TCATCAGGAAGGAAGCAGACAGGGGCCACTCGGAGTCCGCCGAGAGCCACTTGGAAAGGGAAC 1140  
 DB 1081 TCATCAGGAAGGAAGCAGACAGGGGCCACTCGGAGTCCGCCGAGAGCCACTTGGAAAGGGAAC 1140

QY 1141 ACCTGCTCTCCAGATGCACAGCTGGGCTCCGCTGGAAGGTGCTGCACAATGATCTCCGAC 1200  
 DB 1141 ACCTGCTCTCCAGATGCACAGCTGGGCTCCGCTGGAAGGTGCTGCACAATGATCTCCGAC 1200

QY 1201 ATCTCTTCTCTGTAGTGGCTGGGCTCTACAGCCGAGCAGATGCCCCCTCGGAGC 1260  
 DB 1201 ATCTCTTCTCTGTAGTGGGCTCTCTACAGCCGAGCAGATGCCCCCTCGGAGC 1260

QY 1261 CTGGAAACGGGACAGGACAGAGCACAGCTTCTCCCGGATCTCGGACTCTCGGTACAGT 1320  
 DB 1261 CTGGAAACGGGACAGGACAGGACAGGACAGCTTCTCCCGGATCTCGGACTCTCGGTACAGT 1320

QY 1321 GTGGAAAGCCAGCTCTCCAGGCCACGAAAGTCTCTGAGCAGCCTGTATCTTCTGCTCA 1380  
 DB 1321 GTGGAAAGCCAGCTCTCCAGGCCACGAAAGTCTCTGAGCAGCCTGTATCTTCTGCTCA 1380

QY 1381 GTGCCAGAGTCCATGACAATTAGTGAACCTGCGCCAGGCCACTGTGCCATGATGAACAG 1440  
 DB 1381 GTGCCAGAGTCCATGACAATTAGTGAACCTGCGCCAGGCCACTGTGCCATGATGAACAG 1440

QY 1441 AAGGATGAGCTGGAGGAGGAGAAACAGATCACTCGGAAACCTGCTCGACGCTGAGATGGAG 1500  
 DB 1441 AAGGATGAGCTGGAGGAGGAGAAACAGATCACTCGGAAACCTGCTCGACGCTGAGATGGAG 1500

QY 1501 CACTAGCCCGCTCCGGCAAGAGGTGAGACCTTGAAGAAAGGAGTGGCTGAAACAGGAG 1560  
 DB 1501 CACTAGCCCGCTCCGGCAAGAGGTGAGACCTTGAAGAAAGGAGTGGCTGAAACAGGAG 1560

QY 1561 GAGCGCAGGGCATGAAGTCCAGGCGCTGGCCAGCTATCTTTGCTATTTTGTGAGGAGA 1620  
 DB 1561 GAGCGCAGGGCATGAAGTCCAGGCGCTGGCCAGCTATCTTTGCTATTTTGTGAGGAGA 1620

QY 1621 TTCTAAACCCACGTGAGAACCATGTGGTGGAGAAATGGAGGGGAGAGAGAAATCCAAACAGT 1680  
 DB 1621 TTCTAAACCCACGTGAGAACCATGTGGTGGAGAAATGGAGGGGAGAGAGAAATCCAAACAGT 1680

QY 1681 TCTGTAGTCTCATTTGAGTCTCTGGATCCAGTCTTTCTCTGAGCTGTGTTTCTCTCTGG 1740  
 DB 1681 TCTGTAGTCTCATTTGAGTCTCTGGATCCAGTCTTTCTCTGAGCTGTGTTTCTCTCTGG 1740



QY 1081 TCATCAGGAAGGAGACACAGGGGCCACTCGAGTCGCCCGAGAGCCACTGGAGGGAAC 1140  
 Db 1196 TCATCAGGAAGGAGACACAGGGGCCACTCGAGTCGCCCGAGAGCCACTGGAGGGAAC 1255  
 QY 1141 ACCTGCTCTCCAGATGACACAGCTGGGCTCGCTGAAAGTGTGCACAAATGACTCCGAC 1200  
 Db 1256 ACCTGCTCTCCAGATGACACAGCTGGGCTCGCTGAAAGTGTGCACAAATGACTCCGAC 1315  
 QY 1201 ATCTCTTCTCCCTAGTGGCTGCTCTACAGCCGAGAGATGCCCCCTCGAAGC 1260  
 Db 1316 ATCTCTTCTCCCTAGTGGCTGCTCTACAGCCGAGAGATGCCCCCTCGAAGC 1375  
 QY 1261 CTGGAGACGGGACAGACAGAGGACACGTTCTCCCGATCTCGGACTTCGGTACAGT 1320  
 Db 1376 CTGGAGACGGGACAGACAGAGGACACGTTCTCCCGATCTCGGACTTCGGTACAGT 1435  
 QY 1321 GTGGAAGCCAGCTCTCCAGGCCACGGAAGTCTCTGAGCAGCTGTACCTTCTGCTCA 1380  
 Db 1436 GTGGAAGCCAGCTCTCCAGGCCACGGAAGTCTCTGAGCAGCTGTACCTTCTGCTCA 1494  
 QY 1381 GTGCCAGAGTCCATGACAAATTAGTGAATCGCCGAGGCCACTGTGCCATGATGAACAG 1440  
 Db 1495 GTGCCAGAGTCCATGACAAATTAGTGAATCGCCGAGGCCACTGTGCCATGATGAACAG 1554  
 QY 1441 AAGGATGAGCTGGAGGAGGAGACAGATCACTGCGAACTGCTGACCGTGGATGGAG 1500  
 Db 1555 AAGGATGAGCTGGAGGAGGAGACAGATCACTGCGAACTGCTGACCGTGGATGGAG 1614  
 QY 1501 CACTCAGCCGCTCCGGCAAGAGGTGGACACCTTGAAAAGGAGGTGGCTGAACAGGAG 1560  
 Db 1615 CACTCAGCCGCTCCGGCAAGAGGTGGACACCTTGAAAAGGAGGTGGCTGAACAGGAG 1674  
 QY 1561 GAGCGCAGGCGATGAAGTCCAGGCGCTGGCCAGTATCTTTGCTATTTTGTGAGGAGA 1620  
 Db 1675 GAGCGCAGGCGATGAAGTCCAGGCGCTGGCCAGTATCTTTGCTATTTTGTGAGGAGA 1734  
 QY 1621 TTCTACCCACAGTGAGACCATGCTGGTGAGAAATGGAGGAGAGAAATCCACAGT 1680  
 Db 1735 TTCTACCCACAGTGAGACCATGCTGGTGAGAAATGGAGGAGAGAAATCCACAGT 1794  
 QY 1681 TCCTGATAGTCTCATTTGAGCTCTCGATCCAGTCTTTCTGAAGCTGTGTTTCTCTGG 1740  
 Db 1795 TCCTGATAGTCTCATTTGAGCTCTCGATCCAGTCTTTCTGAAGCTGTGTTTCTCTGG 1854  
 QY 1741 ACTTTTCATGATGTGAGGCAATAAATTTGCTTCATTTCTCTTG 1782  
 Db 1855 ACTTTTCATGATGTGAGGCAATAAATTTGCTTCATTTCTCTTG 1896

## RESULT 3

AA004335  
 ID AAD04335 standard; cDNA; 2873 BP.

XX AC AAD04335;

XX DT 04-JUL-2001 (first entry)

XX DE Human TNF receptor-associated factor binding protein, IREN-E cDNA.

XX KW Human; Tumour Necrosis Factor; TNF; TNF Receptor Associated Factor;  
 KW TRAF2; TRAF2 binding protein; IREN; IkappaB Regulator; IREN-E;  
 KW immunosuppressive; nuclear factor-kappaB; NF-kappaB; cytostatic; tumour;  
 KW AIDS; acquired immune deficiency syndrome; rheumatic disease; apoptosis;  
 KW autoimmune disease; septic shock; graft-vs-host reaction; inflammation;  
 KW anorexia; anti-HIV; therapy; ss.

OS Homo sapiens.

XX Key Location/Qualifiers  
 PH CDS 1..2355

FT /\*tag= a

FT /product= "Human TRAF2-binding protein isoform, IREN-E"

XX PN WO2001163114-A1.  
 XX PD 08-MAR-2001.  
 XX PF 31-AUG-2000; 2000WO-IL000517.  
 XX PR 02-SEP-1999; 99IL-00131719.  
 XX PA (YEDA ) YEDA RES & DEV CO LTD.  
 XX PI Wallach D, Malinin N, Sinha I, Leu S;  
 XX WPI: 2001-281387/29.  
 XX P-PSDB; AAE00685.  
 PT New DNA sequence encoding Tumor Necrosis Factor receptor associated  
 PT factor (TRAF) binding proteins (IREN) for treatment or prevention of  
 PT pathological conditions associated with NF-kappaB induction.  
 XX Claim 1; Fig 5; 118pp; English.  
 CC The present cDNA sequence encodes human tumour necrosis factor (TNF)  
 CC receptor-associated factor (TRAF2) binding protein isoform, IREN-E. A  
 CC fragment of this IREN-E molecule is capable of binding to TRAF2 protein  
 CC at position 225-501. The invention relates to human tumour necrosis  
 CC factor (TNF) receptor-associated factor (TRAF2) binding protein  
 CC designated as IREN (IkappaB Regulator), its isoforms IREN-10B, IREN-E and  
 CC their corresponding cDNA molecules. IREN is useful for  
 CC modulating/mediating the activity of transcription factor NF (Nuclear  
 CC Factor)-kappaB or any other intracellular signalling activity mediated by  
 CC TRAF2. IREN is useful in the prevention and treatment of a pathological  
 CC condition associated with NF-kappaB induction (abnormal) e.g. AIDS  
 CC (acquired immune deficiency syndrome), autoimmune diseases, tumours,  
 CC rheumatic diseases, anorexia, septic shock and graft-vs-host reactions.  
 CC IREN also plays an important role in the control of inflammation and  
 CC other non-apoptotic effects of TNF as well as in the control of  
 CC apoptosis. The invention also relates to method for screening,  
 CC identifying and producing a molecule capable of modulating activities  
 CC mediated by IREN. IREN antibodies are useful for the purification of new  
 CC proteins from different sources, including cell extracts or transformed  
 CC cell lines, in addition IREN can be used in diagnostic purposes for  
 CC identifying disorders related to abnormal functioning of cellular effects  
 CC mediated directly by TRAF proteins

SQ Sequence 2873 BP; 756 A; 736 C; 806 G; 575 T; 0 U; 0 Other;

Query Match 89.5%; Score 1595; DB 4; Length 2873;

Best Local Similarity 100.0%; Pred. No. 0;

Matches 1595; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 ATGAGCGGATCAGAAACAATGACAAAACAAATTTCTGTCGAGCGACTGCTGGATGCA 60

Db 1 ATGAGCGGATCAGAAACAATGACAAAACAAATTTCTGTCGAGCGACTGCTGGATGCA 60

QY 61 GTCAAAACAGTGCAGATCCGCTTTGAGGGGAGAAAGAGATTGCTCCGATCCGACAGC 120

Db 61 GTCAAAACAGTGCAGATCCGCTTTGAGGGGAGAAAGAGATTGCTCCGATCCGACAGC 120

QY 121 AGGGTCACCTGTCTGTGTGCCAGTTTGAAGCCGCTCTGCAGCATGGCTTGAAGAGAGT 180

Db 121 AGGGTCACCTGTCTGTGTGCCAGTTTGAAGCCGCTCTGCAGCATGGCTTGAAGAGAGT 180

QY 181 CGAGGATTTGGACTCAGCGGGCAGCGATCAAGCAGGACGCGGCTTTGCCAGCAAAACC 240

Db 181 CGAGGATTTGGACTCAGCGGGCAGCGATCAAGCAGGACGCGGCTTTGCCAGCAAAACC 240

QY 241 GAAACAGAGCCGCTGTCTGTGGTACTACGTGAAGAGGCTCTCAACAGCACGAGCTGCAG 300

Db 241 GAAACAGAGCCGCTGTCTGTGGTACTACGTGAAGAGGCTCTCAACAGCACGAGCTGCAG 300

QY 301 CGCTTCTACTTCCTGCGCCACATTCGCTCAGACGTGGCGCGGGTTCGCGCTGCGTGGC 360

Db 301 CGCTTCTACTTCCTGCGCCACATTCGCTCAGACGTGGCGCGGGTTCGCGCTGCGTGGC 360

Db 301 CGTTTCTACTCCCTCGCGGCCACATCCCTCAGACGTGGGCGGGGTCGCGCCTGGCTGCGC 360  
Qy 361 TGTGCGCTCAACGAACACTCCCTCGAGCGCTACCTGCACATGCTCTCTGCGCGACCGCTGC 420  
Db 361 TGTGCGCTCAACGAACACTCCCTCGAGCGCTACCTGCACATGCTCTCTGCGCGACCGCTGC 420  
Qy 421 AGGCTGAGCACTTTTATGAAGACTGTGCTTTTGTGTGATGATGAAGAAGGTCCAGTATG 480  
Db 421 AGGCTGAGCACTTTTATGAAGACTGTGCTTTTGTGTGATGATGAAGAAGGTCCAGTATG 480  
Qy 481 CTTCTACATGCGACAGGCTGAACCTCCATCTCTTTTGGATTTAAACATCGACAAACAG 540  
Db 481 CTTCTACATGCGACAGGCTGAACCTCCATCTCTTTGGATTTAAACATCGACAAACAG 540  
Qy 541 GATTTGAAACGGGACAGTAAAGTTTGTCTCCACCGTTTCAGACCTCTTAAAGAGTCAACG 600  
Db 541 GATTTGAAACGGGACAGTAAAGTTTGTCTCCACCGTTTCAGACCTCTTAAAGAGTCAACG 600  
Qy 601 CAGAACGTGACTCTTCTGCTGAAGAGTCCACGACGAGGAGTGAAGAGTCTGTTAGGGAG 660  
Db 601 CAGAACGTGACTCTTCTGCTGAAGAGTCCACGACGAGGAGTGAAGAGTCTGTTAGGGAG 660  
Qy 661 ATCAGAGCTCTCTGCGCTCTCCATCTCTCAACCTGAACAGGACGACCCCTTG 720  
Db 661 ATCAGAGCTCTCTGCGCTCTCCATCTCTCAACCTGAACAGGACGACCCCTTG 720  
Qy 721 CCTGCTGTCCAGGAATGTGCTGCTGATGCCAAATGCAAAAAGGAGCGGAAAGAAA 780  
Db 721 CCTGCTGTCCAGGAATGTGCTGCTGATGCCAAATGCAAAAAGGAGCGGAAAGAAA 780  
Qy 781 AAGAAAGTGACCAATATCTCATTTGATGATGAAGAGATGAGAGAACTCTGGGGAC 840  
Db 781 AAGAAAGTGACCAATATCTCATTTGATGATGAAGAGATGAGAGAACTCTGGGGAC 840  
Qy 841 GTGTTTAAAAAGACACCTTGGGCGGAGAGCTCAGAGGACAACTCCGACCGCTCTCT 900  
Db 841 GTGTTTAAAAAGACACCTTGGGCGGAGAGCTCAGAGGACAACTCCGACCGCTCTCT 900  
Qy 901 GTCAATATCATGTCCGCTTTTGAAGCCCTTCCGCGCTAACTCCAATGGAAGTCAGAGC 960  
Db 901 GTCAATATCATGTCCGCTTTTGAAGCCCTTCCGCGCTAACTCCAATGGAAGTCAGAGC 960  
Qy 961 AGCAACTCATGGAATAATGATTCCTGTCTTTGAACGGGGAGTTTGGGTACCAAGAGCTT 1020  
Db 961 AGCAACTCATGGAATAATGATTCCTGTCTTTGAACGGGGAGTTTGGGTACCAAGAGCTT 1020  
Qy 1021 GATGTGAAAAGCATCGATGAAGATGTGATGAAAACGAAGATGAGTGTATGGAAAC 1080  
Db 1021 GATGTGAAAAGCATCGATGAAGATGTGATGAAAACGAAGATGAGTGTATGGAAAC 1080  
Qy 1081 TCATCAGGAAGGAAGCACAGGGGCCACTCGGAGTCGCCGAGAACCACTGGAAGGGAAC 1140  
Db 1081 TCATCAGGAAGGAAGCACAGGGGCCACTCGGAGTCGCCGAGAACCACTGGAAGGGAAC 1140  
Qy 1141 ACCTGCTCTCCAGATGCAAGCTGGGCTCCGCTGAAGGTGCTGCAAAATGATCTCGAC 1200  
Db 1141 ACCTGCTCTCCAGATGCAAGCTGGGCTCCGCTGAAGGTGCTGCAAAATGATCTCGAC 1200  
Qy 1201 ATCTCTTCTCCGTGTCAGTGGGCTCTACAGCCAGAGATGCCCTCCGAGC 1260  
Db 1201 ATCTCTTCTCCGTGTCAGTGGGCTCTACAGCCAGAGATGCCCTCCGAGC 1260  
Qy 1261 CTGAGAAACGGGACAGGACCAAGAGACCACTTCTCCGCGATCTCGGACTTCGGTACAGT 1320  
Db 1261 CTGAGAAACGGGACAGGACCAAGAGACCACTTCTCCGCGATCTCGGACTTCGGTACAGT 1320  
Qy 1321 GTGAGAGCCAGCTCTCCAGGCCACGGAAGTCTCTGAGCAGCCTGTACCTTCTGCTCTCA 1380  
Db 1321 GTGAGAGCCAGCTCTCCAGGCCACGGAAGTCTCTGAGCAGCCTGTACCTTCTGCTCTCA 1380  
Qy 1381 GTGCCAGAGTCCATGACAAATGATGAACTGCGCCAGGACCTGTGGCCATGATGAACAGG 1440  
Db 1381 GTGCCAGAGTCCATGACAAATGATGAACTGCGCCAGGACCTGTGGCCATGATGAACAGG 1440

Qy 1441 AAGGATGAGCTGGAGGAGGAACAGATCACTGCGAAACCTGCTCGACGGTGAGATGGAG 1500  
Db 1441 AAGGATGAGCTGGAGGAGGAACAGATCACTGCGAAACCTGCTCGACGGTGAGATGGAG 1500  
Qy 1501 CACTCAGCGCGCTCCGGCAAGAGGTGGACACCTTGAAAAGAGGAGGTGGCTGAACAGAG 1560  
Db 1501 CACTCAGCGCGCTCCGGCAAGAGGTGGACACCTTGAAAAGAGGAGGTGGCTGAACAGAG 1560  
Qy 1561 GAGCGGAGGCGATGAAGGTCCAGGCGCTGGCCAG 1595  
Db 1561 GAGCGGAGGCGATGAAGGTCCAGGCGCTGGCCAG 1595  
RESULT 4  
AAD04334  
ID AAD04334 standard; cDNA; 3139 BP.  
XX  
AC AAD04334;  
XX  
DT 04-JUL-2001 (first entry)  
XX  
DE Human TNF receptor-associated factor binding protein, IREN-10B cDNA.  
XX  
KW Human; Tumour Necrosis Factor; TNF; TNF Receptor Associated Factor;  
TRAF2; TRAF2 binding protein; IkappaB Regulator; IREN-10B;  
immunosuppressive; nuclear factor-kappaB; NF-kappaB; cytostatic; tumour;  
AIDS; acquired immune deficiency syndrome; rheumatic disease; apoptosis;  
autoimmune disease; septic shock; graft-vs-host reaction; inflammation;  
anorexia; anti-HIV; therapy; ss.  
XX  
OS Homo sapiens.  
XX  
FH Key Location/Qualifiers  
CDS 1..2442  
FT /\*tag= a  
FT /product= "Human TRAF2-binding protein isoform, IREN-10B"  
XX  
WO200116314-A1.  
XX  
PN 08-MAR-2001.  
XX  
PP 31-AUG-2000; 2000WO-IL000517.  
XX  
PR 02-SEP-1999; 99IL-00131719.  
XX  
PA (YEDA ) YEDA RES & DEV CO LTD.  
XX  
PI Wallach D, Malinin N, Sinha I, Leu S;  
XX  
WPI; 2001-281387/29.  
P-PSDB; AAE00684.  
XX  
FT New DNA sequence encoding Tumor Necrosis Factor receptor associated  
factor (TRAF) binding proteins (IREN) for treatment or prevention of  
pathological conditions associated with NF-kappaB induction.  
XX  
PS Claim 1; Fig 4; 118pp; English.  
XX  
CC The present cDNA sequence encodes human tumour necrosis factor (TNF)  
receptor-associated factor (TRAF2) binding protein isoform, IREN-10B. A  
fragment of this IREN-10B molecule is capable of binding to TRAF2 protein  
at position 225-501. The invention relates to human tumour necrosis  
factor (TNF) receptor- associated factor (TRAF2) binding protein  
designated as IREN (IkappaB Regulator). IREN is useful for  
modulating/mediating the activity of transcription factor NF (Nuclear  
Factor)-kappaB or any other intracellular signalling activity mediated by  
TRAF2. IREN is useful in the prevention and treatment of a pathological  
condition associated with NF-kappaB induction (abnormal) e.g. AIDS  
(acquired immune deficiency syndrome), autoimmune diseases, tumours,  
rheumatic diseases, anorexia, septic shock and graft-vs-host reactions.  
CC IREN also plays an important role in the control of inflammation and

CC other non-apoptotic effects of TNF as well as in the control of  
CC apoptosis. The invention also relates to method for screening,  
CC identifying and producing a molecule capable of modulating activities  
CC mediated by IREN. IREN antibodies are useful for the purification of new  
CC proteins from different sources, including cell extracts or transformed  
CC cell lines, in addition IREN can be used in diagnostic purposes for  
CC identifying disorders related to abnormal functioning of cellular effects  
CC mediated directly by TRAF proteins

XX SQ Sequence 3139 BP; 797 A; 851 C; 875 G; 616 T; 0 U; 0 Other;

Query Match		89.5%;	Score 1595;	DB 4;	Length 3139;
Best Local Similarity		100.0%;	Pred. No. 0;		
Matches 1595;		Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;
QY	1	ATGAGCGGATCACAGAACATGCAAAAGACAAATTTCTGCTGAGGAGCTGCTGGATGCA	60		
DB	1	ATGAGCGGATCACAGAACATGCAAAAGACAAATTTCTGCTGAGGAGCTGCTGGATGCA	60		
QY	61	GTGAAACAGTCCAGATCCGCTTTGGAGGGAGAAAGAGATTCCCTCGGATTCGGACAGC	120		
DB	61	GTGAAACAGTCCAGATCCGCTTTGGAGGGAGAAAGAGATTCCCTCGGATTCGGACAGC	120		
QY	121	AGGGTCACCTGTCTGTGTCGCCAGTTTGAAGCCGCTCTGCAGCATGCTTGAAGAGGAGT	180		
DB	121	AGGGTCACCTGTCTGTGTCGCCAGTTTGAAGCCGCTCTGCAGCATGCTTGAAGAGGAGT	180		
QY	181	CGAGGATTGGCACTACAGCGGAGCGGATCAAGCAGGAGCGGGCTTTGCGACAAACC	240		
DB	181	CGAGGATTGGCACTACAGCGGAGCGGATCAAGCAGGAGCGGGCTTTGCGACAAACC	240		
QY	241	GAAACAGAGCCGTGTTCTGTTACTAGTGAAGAGGAGTCTCAACAGACAGAGCTGCAG	300		
DB	241	GAAACAGAGCCGTGTTCTGTTACTAGTGAAGAGGAGTCTCAACAGACAGAGCTGCAG	300		
QY	301	CGCTTCTACTCCCTGGCCACATCGCTCAGAGTGGCGGGGTCGCGCTCGCTCGCG	360		
DB	301	CGCTTCTACTCCCTGGCCACATCGCTCAGAGTGGCGGGGTCGCGCTCGCTCGCG	360		
QY	361	TGTGCCCTCAACGAACACTCCCTGGAGCGCTACCTGCACATGCTCTCTGGCCGCGCTGC	420		
DB	361	TGTGCCCTCAACGAACACTCCCTGGAGCGCTACCTGCACATGCTCTCTGGCCGCGCTGC	420		
QY	421	AGCTGAGCACTTTTATGAAGACTGCTCTTTTGTGATGATGAAGAGGTCAGTATG	480		
DB	421	AGCTGAGCACTTTTATGAAGACTGCTCTTTTGTGATGATGAAGAGGTCAGTATG	480		
QY	481	CTTCTTACCATGGCAGAGGCTGAACTCCATCTCTTTCCGATTAAACATCGACAACAG	540		
DB	481	CTTCTTACCATGGCAGAGGCTGAACTCCATCTCTTTCCGATTAAACATCGACAACAG	540		
QY	541	GATTGAAACGGGAGAGTAAATTTGCTCCACCGTTTTCAGACTCTTTAAAGAGTCAACG	600		
DB	541	GATTGAAACGGGAGAGTAAATTTGCTCCACCGTTTTCAGACTCTTTAAAGAGTCAACG	600		
QY	601	CAGAACGTGACCTCTTCTGAGGAGTCCACCGAGAGTGGAGCTGTTTCAGGGAG	660		
DB	601	CAGAACGTGACCTCTTCTGAGGAGTCCACCGAGAGTGGAGCTGTTTCAGGGAG	660		
QY	661	ATCACAGCTCTCTGCGCTCTCCATCTCATCAAACTCTGAAACAGGAGACCGCCCTTG	720		
DB	661	ATCACAGCTCTCTGCGCTCTCCATCTCATCAAACTCTGAAACAGGAGACCGCCCTTG	720		
QY	721	CTGTGTGTCAGGAATGTCAGTCTGATGTCGCAATGCAAAAGAGGAGCGGAAGAAA	780		
DB	721	CTGTGTGTCAGGAATGTCAGTCTGATGTCGCAATGCAAAAGAGGAGCGGAAGAAA	780		
QY	781	AAGAAAGTGAACCAATCTCAATTTGATGAGGAGGAGTGAAGAGTCTGCGGAC	840		
DB	781	AAGAAAGTGAACCAATCTCAATTTGATGAGGAGGAGTGAAGAGTCTGCGGAC	840		
QY	841	GTGTTTAAAGACACCTGGGCGGAGGAGCTCAGAGGACAACTCCGACCGCTCTCT	900		

DB	841	GTGTTTAAAGACACCTGGGCGAGGGAGAGCTCAGAGGACAACTCCGACCGCTCTCT	900
QY	901	GTCAATATCATGTCGCGCTTTTGAAGCCCTTTCGGGCTTAACCTCCAATGGAAGTCAGAGC	960
DB	901	GTCAATATCATGTCGCGCTTTTGAAGCCCTTTCGGGCTTAACCTCCAATGGAAGTCAGAGC	960
QY	961	AGCAACTCATGAAATATGATTCCTCTGCTTTTGAACGGGAGTTTGGGTACCAAGACCTT	1020
DB	961	AGCAACTCATGAAATATGATTCCTCTGCTTTTGAACGGGAGTTTGGGTACCAAGACCTT	1020
QY	1021	GATGTGAAAGCATCGATGATGGAATGAAACCAAGATGACCTGTATGGAAC	1080
DB	1021	GATGTGAAAGCATCGATGATGGAATGAAACCAAGATGACCTGTATGGAAC	1080
QY	1081	TATCATGGAAGGAAGCACAGGGCCACTCGGAGTCGCCCGAGAGCCACTGGAAGGGAAC	1140
DB	1081	TATCATGGAAGGAAGCACAGGGCCACTCGGAGTCGCCCGAGAGCCACTGGAAGGGAAC	1140
QY	1141	ACCTGCTCTCCAGATGCACAGCTGGCTCCGCTGAAGGTGCTGCACATGACTCCGAC	1200
DB	1141	ACCTGCTCTCCAGATGCACAGCTGGCTCCGCTGAAGGTGCTGCACATGACTCCGAC	1200
QY	1201	ATCCTCTTCCCTGTCACTGGCTGCTTACAGCCACAGCAGATGCCCCCTCGGAAGC	1260
DB	1201	ATCCTCTTCCCTGTCACTGGCTGCTTACAGCCACAGCAGATGCCCCCTCGGAAGC	1260
QY	1261	CTGGAGAACGGGACAGGACACAGGACCAAGCTTCTCCCGATCTCTGGACTTCGGTACAGT	1320
DB	1261	CTGGAGAACGGGACAGGACACAGGACCAAGCTTCTCCCGATCTCTGGACTTCGGTACAGT	1320
QY	1321	GTGGAGAACGAGCTCTCCAGGCCACGAACTCTCTGAGCAGCTGTTACCTTCTGCTCA	1380
DB	1321	GTGGAGAACGAGCTCTCCAGGCCACGAACTCTCTGAGCAGCTGTTACCTTCTGCTCA	1380
QY	1381	GTGCCAGAGTCCATGAACTTAGTGAATTCGCGCAGGCCACTGTGCGCATGATGAACAGG	1440
DB	1381	GTGCCAGAGTCCATGAACTTAGTGAATTCGCGCAGGCCACTGTGCGCATGATGAACAGG	1440
QY	1441	AAGGATGAGCTGGAGGAGGAGAACAGATCACTGCGAAACCTGCTCGACGGTGAAGTGAG	1500
DB	1441	AAGGATGAGCTGGAGGAGGAGAACAGATCACTGCGAAACCTGCTCGACGGTGAAGTGAG	1500
QY	1501	CATCAGCGCGCTCCGCGAAGAGGTGACACCTCTGAAAGGAAGGTGCTGAAACAGGAG	1560
DB	1501	CATCAGCGCGCTCCGCGAAGAGGTGACACCTCTGAAAGGAAGGTGCTGAAACAGGAG	1560
QY	1561	GAGCGCAGGCGATGAAGGTCCAGGCGCTGGCCAG	1595
DB	1561	GAGCGCAGGCGATGAAGGTCCAGGCGCTGGCCAG	1595

RESULT 5  
AAH15665  
ID AAH15665 standard; cDNA; 2248 BP.  
XX  
AC AAH15665;  
XX  
XX 26-JUN-2001 (first entry)  
XX  
DE Human cDNA sequence SRQ ID NO:14016.  
XX  
KW Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.  
XX Homo sapiens.  
XX  
XX EP1074617-A2.  
XX  
PD 07-FEB-2001.  
XX  
XX 28-JUL-2000; 2000EP-00116126.  
PF  
XX 29-JUL-1999; 99JP-00248036.  
PR 27-AUG-1999; 99JP-00300253.

PR 11-JAN-2000; 2000JP-00118776.  
PR 02-MAY-2000; 2000JP-00183767.  
PR 09-JUN-2000; 2000JP-00241899.  
XX (HELI-) HELIX RES INST.  
XX  
PI Ota T, Isegai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;  
PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;  
XX WPI; 2001-318749/34.  
XX  
XX Primer sets for synthesizing polynucleotides, particularly the 5602 full-  
PT length cDNAs defined in the specification, and for the detection and/or  
PT diagnosis of the abnormality of the proteins encoded by the full-length  
PT cDNAs.  
XX  
PS Claim 8; SEQ ID NO 14016; 2537pp + Sequence Listing; English.  
XX  
XX The present invention describes primer sets for synthesizing 5602 full-  
CC length cDNAs defined in the specification. Where a primer set comprises:  
CC (a) an oligo-dT primer and an oligonucleotide complementary to the  
CC complementary strand of a polynucleotide which comprises one of the 5602  
CC nucleotide sequences defined in the specification, where the  
CC oligonucleotide comprises at least 15 nucleotides; or (b) a combination  
CC of an oligonucleotide comprising a sequence complementary to the  
CC complementary strand of a polynucleotide which comprises a 5'-end  
CC sequence and an oligonucleotide comprising a sequence complementary to a  
CC polynucleotide which comprises a 3'-end sequence, where the  
CC oligonucleotide comprises at least 15 nucleotides and the combination of  
CC the 5'-end sequence/3'-end sequence is selected from those defined in the  
CC specification. The primer sets can be used in antisense therapy and in  
CC gene therapy. The primers are useful for synthesizing polynucleotides,  
CC particularly full-length cDNAs. The primers are also useful for the  
CC detection and/or diagnosis of the abnormality of the proteins encoded by  
CC the full-length cDNAs. The primers allow obtaining of the full-length  
CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and  
CC AAH13633 to AAH18742 represent human cDNA sequences; AAH92446 to AAH95893  
CC represent human amino acid sequences; and AAH13629 to AAH13632 represent  
CC oligonucleotides, all of which are used in the exemplification of the  
XX present invention  
XX  
SQ Sequence 2248 BP; 548 A; 548 C; 618 G; 534 T; 0 U; 0 Other;

Query Match 63.4%; Score 1129.8; DB 4; Length 2248;  
Best Local Similarity 95.3%; Pred. No. 1.1e-297;  
Matches 1164; Conservative 0; Mismatches 57; Indels 0; Gaps 0;

Qy 1 ATGAGCGGATCACAGAACAAATGACAAAAGACAATTTCTGCTGGAGCGACTCTGGATGCA 60  
Db 63 ATGAGCGGATCACAGAACAAATGACAAAAGACAATTTCTGCTGGAGCGACTCTGGATGCA 122  
Qy 61 GTGAAACAGTCCAGATCCGCTTTGGAGGAGAAAGAGATTTGCTCGGATTCGACAGC 120  
Db 123 GTGAAACAGTCCAGATCCGCTTTGGAGGAGAAAGAGATTTGCTCGGATTCGACAGC 182  
Qy 121 AGGTTCACCTGTCTGTGTGCCAGTTTGAAGCCGTCCTGACAGCTGGCTTGAAGAGGAGT 180  
Db 183 AGGTTCACCTGTCTGTGTGCCAGTTTGAAGCCGTCCTGACAGCTGGCTTGAAGAGGAGT 242  
Qy 181 CGAGGATGGCACTCACAGCGCGCGCGATCAAGCAGGCGAGGGCTTTGCCAGCAAAACC 240  
Db 243 CGAGGATGGCACTCACAGCGCGCGCGATCAAGCAGGCGAGGGCTTTGCCAGCAAAACC 302  
Qy 241 GAAACAGAGCCGCTGTCTGTGTACTACGTGAAGAGGTCCTCAACAGCAGAGCTGCAG 300  
Db 303 GAAACAGAGCCGCTGTCTGTGTACTACGTGAAGAGGTCCTCAACAGCAGAGCTGCAG 362  
Qy 301 CGCTTCTACTCCCTCGCCACATCGCTTCAGACGTGGCGCGGGTCGCGCTTGGCTGCGC 360  
Db 363 CGCTTCTACTCCCTCGCCACATCGCTTCAGACGTGGCGCGGGTCGCGCTTGGCTGCGC 422  
Qy 361 TGTGCCCTCAAGCAACTCCCTGGAGCGCTACCTGCACATGCTCTCTGGCCGACCGCTGC 420

Db 423 TGTGCCCTCAAGCAACACTCCCTGGAGCGCTACCTGCACATGCTCTCTGGCCGACCGCTGC 482  
Qy 421 AGGTTCAGCACTTTTATGAAGACTGGTCTTTTGTGATGGATGAAGAAGGTCACAGTATG 480  
Db 483 AGGTTCAGCACTTTTATGAAGACTGGTCTTTTGTGATGGATGAAGAAGGTCACAGTATG 542  
Qy 481 CTTCTTACCATTGGCAGCAGGTCTGAATCTCATATCTTTTGGCATTAACATCGACAAACAG 540  
Db 543 CTTCTTACCATTGGCAGCAGGTCTGAATCTCATATCTTTTGGCATTAACATCGACAAACAG 602  
Qy 541 GATTGACAGCGGCGAGAGTTAGTTTCTCCCAACCGTTTTCAGACCTCTTAAAGGAGTCAAG 600  
Db 603 GATTGACAGCGGCGAGAGTTAGTTTCTCCCAACCGTTTTCAGACCTCTTAAAGGAGTCAAG 662  
Qy 601 CAGAACGTGACCTCTCTTGTCTGAAGAGTCCACGCAAGGAGTGAGCAGCTGTTCAGGGAG 660  
Db 663 CAGAACGTGACCTCTCTTGTCTGAAGAGTCCACGCAAGGAGTGAGCAGCTGTTCAGGGAG 722  
Qy 661 ATCAAGAGCTCTCTTGGCGTCTCCATCTCATATCAAACTGAAACGAGGAGACCGACCCCTTG 720  
Db 723 ATCAAGAGCTCTCTTGGCGTCTCCATCTCATCAAACTGAAACGAGGAGACCGACCCCTTG 782  
Qy 721 CCTCTCTGTCAGGAAATGTCTGCTGATGCCAAATGCAAAAGGAGCGGAAGAGAAA 780  
Db 783 CCTCTCTGTCAGGAAATGTCTGCTGATGCCAAATGCAAAAGGAGCGGAAGAGAAAG 842  
Qy 781 AAGAAAGTGACCAACAATAATCTCATTTGATGAGGAAGATGAGCAAACTCTGGGGAC 840  
Db 843 AAGAAAGTGACCAACAATAATCTCATTTGATGAGGAAGATGAGCAAACTCTGGGGAC 902  
Qy 841 GTGTTTAAAAAGACACCTGGGGCAGGGGAGAGCTCAGAGGACAACTCCGACCGCTCTCT 900  
Db 903 GTGTTTAAAAAGACACCTGGGGCAGGGGAGAGCTCAGAGGACAACTCCGACCGCTCTCT 962  
Qy 901 GTCAATATCATGTCCGCTTTGAAGCCCTTCGGGCTTAATCTCAATGGAAGTCAAGAC 960  
Db 963 GTCAATATCATGTCCGCTTTGAAGCCCTTCGGGCTTAATCTCAATGGAAGTCAAGAC 1022  
Qy 961 AGCAACTCATGGAATAATGATTCCTCTGTCTTGAACGGGGAGTTTGGGTACCAAGAGCTT 1020  
Db 1023 AGCAACTCATGGAATAATGATTCCTCTGTCTTGAACGGGGAGTTTGGGTACCAAGAGCTT 1082  
Qy 1021 GATGTGAAAGCATCGATGATGAAGATGTGATGAAACGAAAGATGATGATGGAAC 1080  
Db 1083 GATGTGAAAGCATCGATGATGAAGATGTGATGAAACGAAAGATGATGATGGAAC 1142  
Qy 1081 TCATCAGGAAGAGACACGGGCGCACTCGAGTTCGCCCGAGAGCCACTGGAAGGGAGAC 1140  
Db 1143 TCATCAGGAAGAGACACGGGCGCACTCGAGTTCGCCCGAGAGAGTAAAGTTTGTGTGTA 1202  
Qy 1141 ACCTGCTCTCCAGATGACACAGCTGGGCTCCGCTGAAGGTGCTGCACAAATGACTCCGAC 1200  
Db 1203 GGTGAGTCTCACCGTCCCGCAGGCTGGAGTGGCGTGGATCTCAGCTCACTGCAAC 1262  
Qy 1201 ATCTCTTTCCTGTCACTGGC 1221  
Db 1263 CTCACCTCCCGGTTTCAGGC 1283

RESULT 6  
AAH18235  
ID AAH18235 standard; cDNA; 6045 BP.  
XX  
AC AAH18235;  
XX  
XX 26-JUN-2001 (first entry)  
XX  
DE Human cDNA sequence SEQ ID NO:18174.  
XX  
KW Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.  
XX  
XX Homo sapiens.  
XX



PN EP1074617-A2.  
 XX 07-FEB-2001.  
 XX 28-JUL-2000; 2000EP-00116126.  
 XX 29-JUL-1999; 99JP-00248036.  
 PR 27-AUG-1999; 99JP-00300253.  
 PR 11-JAN-2000; 2000JP-00118776.  
 PR 02-MAY-2000; 2000JP-00183767.  
 PR 09-JUN-2000; 2000JP-00241899.  
 XX (HELI-) HELIX RES INST.  
 XX Ota T, Isogai T, Nishikawa T, Hayaishi K, Saito K, Yamamoto J;  
 PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;  
 XX WPI; 2001-318749/34.  
 XX  
 PT Primer sets for synthesizing polynucleotides, particularly the 5602 full-length cDNAs defined in the specification, and for the detection and/or diagnosis of the abnormality of the proteins encoded by the full-length cDNA.  
 PT  
 PT  
 XX Claim 8; SEQ ID NO 18174; 2537pp + Sequence Listing; English.  
 XX  
 CC The present invention describes primer sets for synthesizing 5602 full-length cDNAs defined in the specification. Where a primer set comprises:  
 CC (a) an oligo-dT primer and an oligonucleotide complementary to the  
 CC complementary strand of a polynucleotide which comprises one of the 5602  
 CC nucleotide sequences defined in the specification, where the  
 CC oligonucleotide comprises at least 15 nucleotides; or (b) a combination  
 CC of an oligonucleotide comprising a sequence complementary to the  
 CC complementary strand of a polynucleotide which comprises a 5'-end  
 CC sequence and an oligonucleotide comprising a sequence complementary to a  
 CC polynucleotide which comprises a 3'-end sequence, where the  
 CC oligonucleotide comprises at least 15 nucleotides and the combination of  
 CC the 3'-end sequence and the sequence is selected from those defined in the  
 CC specification. The primer sets can be used in antisense therapy and in  
 CC gene therapy. The primers are useful for synthesizing polynucleotides,  
 CC particularly full-length cDNAs. The primers are also useful for the  
 CC detection and/or diagnosis of the abnormality of the proteins encoded by  
 CC the full-length cDNAs. The primers allow obtaining of the full-length  
 CC cDNAs easily without any specialised methods. AAH03166 to AAH13628 and  
 CC AAH13633 to AAH18742 represent human cDNA sequences; AAB92446 to AAB95893  
 CC represent human amino acid sequences; and AAH13629 to AAH13632 represent  
 CC oligonucleotides, all of which are used in the exemplification of the  
 CC present invention  
 XX  
 SQ Sequence 6045 BP; 1561 A; 1493 C; 1427 G; 1564 T; 0 U; 0 Other;

Query Match 45.8%; Score 816.8; DB 4; Length 6045;  
 Best Local Similarity 97.0%; Pred. No. 6e-212;  
 Matches 854; Conservative 0; Mismatches 22; Indels 4; Gaps 2;

QY 246 AGAGCCCGTGTTCGTACTACGTGAAGAGGCTCCTCAACAGCAGCGAGTGC-AGCGCT 304  
 DB 932 AAAGCCCGTGTTCGTACTACGTGAAGAGGCTCCTCAACAGCAGCGAGTGC-AGCGCT 991  
 QY 305 TCTACTCCCTCGCCACATCGCTCAGACGTGGCGCGGGTCCGCGCTGCTGTG 364  
 DB 992 TCTACTCCCTCGCCACATCGCTCAGACGTGGCGCGGGTCCGCGCTGCTGTG 1051  
 QY 365 CCTCAACGAACACTCCCTCGAGCGCTACTGACATGCTCTCGCGCGAGCCGCTGACGC 424  
 DB 1052 CCTCAACGAACACTCCCTCGAGCGCTACTGACATGCTCTCGCGCGAGCCGCTGACGC 1111  
 QY 425 TGAGCACTTTTATGACACTGCTCTTCTGATGGATGAAGAAGTCCAGTATGCTTC 484  
 DB 1112 TCAGTACTTTTATGACACTGCTCTTCTGATGGATGAAGAAGTCCAGTATGCTTC 1171  
 QY 485 CTACCATGGCAGCGGTCTCTGAACCTCCACTCTCTTGGATTAAACATCGACAACAGGATT 544

DB 1172 CTACCATGGCAGCGGTCCGAACTCCATACTCTTTGCGATTAACTTGAACAACAGGATT 1231  
 QY 545 TGAACGGGCGAGTAAGTTTGTCTCCACCGTTTTCAGACCTCTTTAAAGGAGTCAACGCAGA 604  
 DB 1232 TGAACGGGCGAGTAAGTTTGTCTCCACCGTTTTCAGACCTCTTTAAAGGAGTCAACGCAGA 1291  
 QY 605 AGCTGACCTCCCTTGTCTGAAGGAGTCCACGCAAGGAGTGAAGAGTCTTTCAGGAGATCA 664  
 DB 1292 ATGTGA---CCTTGTCTGAAGGAGTCCACCAAGGAGTGAAGAGTCTTTCAGGAGATCA 1348  
 QY 665 CAGCCTCTCTCTGCCGTCTCCATCTCTCATCAAACTGAAACAGGAGAGCCGACCTTTCCTG 724  
 DB 1349 CAGCCTCTCTCTGCCATCTCCATCTCTCATCAAACTGAAACAGGAGAGCCGACCTTTCCTG 1408  
 QY 725 TCGTGTCCAGGAATGTCTAGTCTGATGCTCAAAATGCAAAAGGAGCGGAAGAGAAAAAGA 784  
 DB 1409 TCGTGTCCAGGAATGTCTAGTCTGATGCTCAAAATGCAAAAGGAGCGGAAGAGAAAAAGC 1468  
 QY 785 AAGTGACCAACATAATCTCATTTTGATGATGAGGAAGATGAGCAGAACTCTGGGACCTGT 844  
 DB 1469 AAGTGACCAACATAATCTCATTTTGATGATGAGGAAGATGAGCAGAACTCTGGGACATGT 1528  
 QY 845 TTAATAAGACACTGGGGCAGGGAGAGCTCAGAGACAACTCCGACCCCTCTCTGTCTCA 904  
 DB 1529 TTAATAAGACACTGGGGCAGGGAGAGCTCAGAGACAACTCCGACCCCTCTCTGTCTCA 1588  
 QY 905 ATATCATGTCCGCTTTGAAAGCCCTTCGGGCTTAACCTCAATGGAAGTCAAGAGCAGCA 964  
 DB 1589 ATATCATGTCCGCTTTGAAAGCCCTTCGGGCTTAACCTCAATGGAAGTCAAGAGCAGCA 1648  
 QY 965 ACTCATGGAAAAATTGATTCCTCTGTTTGAACGGGAGTTTGGGTACCAAGAGCTTGATG 1024  
 DB 1649 ACTCTGGAAAAATTGATTCCTCTGTTTGAACGGGAGTTTGGGTACCAAGAGCTTGATG 1708  
 QY 1025 TGAAGAAGCATCATGATGAAGATGTGGATGAAAAAGAGATGACGCTGTATGAAACTCAT 1084  
 DB 1709 TGAAGAAGCATCATGATGAAGATGTGGATGAAAAAGAGATGACGCTGTATGAAACTCAT 1768  
 QY 1085 CAGGAAGGAGACAGCGGGCCACTCGGACTCCGCGGAA 1124  
 DB 1769 CAGGAAGGAGACAGCGGGCCACTCAGAGTCCGCGGAGAA 1808  
 XX  
 XX AAH04973;  
 XX  
 DT 26-JUN-2001 (first entry)  
 XX  
 DE Human cDNA clone (5'-primer) SEQ ID NO:1808.  
 XX  
 KW Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.  
 XX  
 OS Homo sapiens.  
 XX  
 PN EP1074617-A2.  
 XX  
 PD 07-FEB-2001.  
 XX  
 PF 28-JUL-2000; 2000EP-00116126.  
 XX  
 PR 29-JUL-1999; 99JP-00248036.  
 PR 27-AUG-1999; 99JP-00300253.  
 PR 11-JAN-2000; 2000JP-00118776.  
 PR 02-MAY-2000; 2000JP-00183767.  
 PR 09-JUN-2000; 2000JP-00241899.  
 XX  
 XX (HELI-) HELIX RES INST.  
 XX  
 PI Ota T, Isogai T, Nishikawa T, Hayaishi K, Saito K, Yamamoto J;  
 PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;



XX WPI; 2001-318749/34.  
XX Primer sets for synthesizing polynucleotides, particularly the 5602 full-length cDNAs defined in the specification, and for the detection and/or diagnosis of the abnormality of the proteins encoded by the full-length cDNAs.  
XX Claim 1; SEQ ID NO 1808; 2537pp + Sequence Listing; English.  
XX The present invention describes primer sets for synthesizing 5602 full-length cDNAs defined in the specification. Where a primer set comprises: (a) an oligo-dr primer and an oligonucleotide complementary to the complementary strand of a polynucleotide which comprises one of the 5602 nucleotide sequences defined in the specification, where the oligonucleotide comprises at least 15 nucleotides; or (b) a combination of an oligonucleotide comprising a sequence complementary to the complementary strand of a polynucleotide which comprises a 5'-end sequence and an oligonucleotide comprising a sequence complementary to a polynucleotide which comprises a 3'-end sequence, where the oligonucleotide comprises at least 15 nucleotides and the combination of the 5'-end sequence/3'-end sequence is selected from those defined in the specification. The primer sets can be used in antisense therapy and in gene therapy. The primers are useful for synthesizing polynucleotides, particularly full-length cDNAs. The primers are also useful for the detection and/or diagnosis of the abnormality of the proteins encoded by the full-length cDNAs. The primers allow obtaining of the full-length cDNAs easily without any specialised methods. AAH03166 to AAH13628 and AAH13633 to AAH18742 represent human cDNA sequences; AAH92446 to AAH95893 represent human amino acid sequences; and AAH13629 to AAH13632 represent polynucleotides, all of which are used in the exemplification of the present invention  
XX SQ Sequence 730 BP; 173 A; 200 C; 217 G; 137 T; 0 U; 3 Other;  
Query Match 35.9%; Score 639.8; DB 4; Length 730;  
Best Local Similarity 99.0%; Pred. No. 5e-164;  
Matches 663; Conservative 0; Mismatches 5; Indels 2; Gaps 2;  
QY 1 ATGAGCGGATCACAGAAACAATGACAAAAGACAATTTCTGCTGGAGCGCACTGCTGGATGCA 60  
DB 63 ATGAGCGGATCACAGAAACAATGACAAAAGACAATTTCTGCTGGAGCGCACTGCTGGATGCA 122  
QY 61 GTCAAAACAGTCCAGATCCGCTTTGGAGGAGAAAGAGATGCTCGGATTCGACAGC 120  
DB 123 GTGAAACAGTCCAGATCCGCTTTGGAGGAGAAAGAGATGCTCGGATTCGACAGC 182  
QY 121 AGGTCACCTGTCTGTGTGCCAGTTTGAAGCCGCTCTGAGCATGCTTGAAGAGAGT 180  
DB 183 AGGTCACCTGTCTGTGTGCCAGTTTGAAGCCGCTCTGAGCATGCTTGAAGAGAGT 242  
QY 181 CGAGGATTGGCACTCAGCGGCGCAGCGATCAAGCAGCAGCGGGCTTTGCCAGCAAAACC 240  
DB 243 CGAGGATTGGCACTCAGCGGCGCAGCGATCAAGCAGCAGCGGGCTTTGCCAGCAAAACC 302  
QY 241 GAAACAGAGCCGCTGTTCTGTGTACTACGTGAAGAGATGCTCTCAACAGCAGAGCTGCAG 300  
DB 303 GAAACAGAGCCGCTGTTCTGTGTACTACGTGAAGAGATGCTCTCAACAGCAGAGCTGCAG 362  
QY 301 CGCTTCTACTCCCTGCGCCACATCGCTCAGACGTGGCGGGGTCGGCTGCTGCGC 360  
DB 363 CGCTTCTACTCCCTGCGCCACATCGCTCAGACGTGGCGGGGTCGGCTGCTGCGC 422  
QY 361 TGTGCTCTCAACGAACACTCCCTGGAGCGCTTACCTGCACATGCTCTCTGGCGGCGCGTGC 420  
DB 423 TGTGCTCTCAACGAACACTCCCTGGAGCGCTTACCTGCACATGCTCTCTGGCGGCGCGTGC 482  
QY 421 AGCTGAGCACTTTTATGAAGACTGGTCTTTTGTGATGATGAAGAAAGTCCAGTATG 480  
DB 483 AGCTGAGCACTTTTATGAAGACTGGTCTTTTGTGATGATGAAGAAAGTCCAGTATG 542  
QY 481 CTTCTCTACCATGGCAGAGGTCTGAACCTCCATCTCTTTGGGATTAACATCGACAAACAG 540  
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DB 543 CTTCTCTACCATGGCAGAGGTCTGAACCTCCATCTCTTTGGGATTAACATCGACAAACAG 602  
QY 541 GATTTCGAACGGGAGAGTAAGTTTGTCTCCACCGTTTTCAGACCTCTTTAAAGGAGTCAACG 600  
DB 603 GATTTCGAACGGGAGAGTAAGTTTGTCTCCACCGTTTTCAGACCTCTTTAAAGGAGTCAACG 662  
QY 601 CAGAACGTGACCTCTCTTGTCTGAAGAGTCCACGCAAGAGTGAAGAGTCTGTTTTCAGGGAG 660  
DB 663 CANAACGTGA-CTNCTTGTCTGAAGAGTCCACGCAAGAGTGANCA-CTTGTTCAGAGGAG 720  
QY 661 ATCAGAGCCT 670  
DB 721 ATCAGAGCCT 730  
RESULT 8  
AAK67282  
ID AAK67282 standard; DNA; 33147 BP.  
XX AAK67282;  
AC AAK67282;  
XX DT 06-NOV-2001 (first entry)  
XX Human immune/haematopoietic antigen genomic sequence SEQ ID NO:22094.  
XX Human; immune; haematopoietic; immune/haematopoietic antigen; cancer;  
KW cytostatic; gene therapy; vaccine; metastasis; ds.  
XX Homo sapiens.  
OS Homo sapiens.  
XX WO200157182-A2.  
PN 09-AUG-2001.  
XX PD 17-JAN-2001; 2001WO-US001354.  
XX PF 31-JAN-2000; 2000US-0179065P.  
PR 04-FEB-2000; 2000US-0180628P.  
PR 24-FEB-2000; 2000US-0184664P.  
PR 02-MAR-2000; 2000US-0186350P.  
PR 16-MAR-2000; 2000US-0189874P.  
PR 17-MAR-2000; 2000US-0190076P.  
PR 18-APR-2000; 2000US-0198123P.  
PR 19-MAY-2000; 2000US-0205515P.  
PR 07-JUN-2000; 2000US-0209467P.  
PR 28-JUN-2000; 2000US-0214886P.  
PR 30-JUN-2000; 2000US-0215135P.  
PR 07-JUL-2000; 2000US-0216647P.  
PR 07-JUL-2000; 2000US-0216880P.  
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PR 14-JUL-2000; 2000US-0218290P.  
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PR 14-AUG-2000; 2000US-0225470P.  
PR 14-AUG-2000; 2000US-0225477P.  
PR 14-AUG-2000; 2000US-0225757P.  
PR 14-AUG-2000; 2000US-0225758P.  
PR 14-AUG-2000; 2000US-0225759P.  
PR 18-AUG-2000; 2000US-0226279P.  
PR 22-AUG-2000; 2000US-0226681P.  
PR 22-AUG-2000; 2000US-0226688P.  
PR 23-AUG-2000; 2000US-0227182P.  
PR 23-AUG-2000; 2000US-0227009P.  
PR 30-AUG-2000; 2000US-0228924P.  
PR 01-SEP-2000; 2000US-0229287P.

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PR 01-SEP-2000; 2000US-0229343P.
PR 01-SEP-2000; 2000US-0229344P.
PR 01-SEP-2000; 2000US-0229345P.
PR 05-SEP-2000; 2000US-0229509P.
PR 05-SEP-2000; 2000US-0229513P.
PR 06-SEP-2000; 2000US-0230437P.
PR 06-SEP-2000; 2000US-0230438P.
PR 08-SEP-2000; 2000US-0231242P.
PR 08-SEP-2000; 2000US-0231243P.
PR 08-SEP-2000; 2000US-0231244P.
PR 08-SEP-2000; 2000US-0231413P.
PR 08-SEP-2000; 2000US-0231414P.
PR 08-SEP-2000; 2000US-0232080P.
PR 08-SEP-2000; 2000US-0232081P.
PR 12-SEP-2000; 2000US-0231968P.
PR 14-SEP-2000; 2000US-0232397P.
PR 14-SEP-2000; 2000US-0232398P.
PR 14-SEP-2000; 2000US-0232399P.
PR 14-SEP-2000; 2000US-0232400P.
PR 14-SEP-2000; 2000US-0232401P.
PR 14-SEP-2000; 2000US-0233063P.
PR 14-SEP-2000; 2000US-0233064P.
PR 14-SEP-2000; 2000US-0233065P.
PR 21-SEP-2000; 2000US-0234223P.
PR 21-SEP-2000; 2000US-0234274P.
PR 25-SEP-2000; 2000US-0234997P.
PR 25-SEP-2000; 2000US-0234998P.
PR 26-SEP-2000; 2000US-0235484P.
PR 27-SEP-2000; 2000US-0235834P.
PR 27-SEP-2000; 2000US-0235836P.
PR 29-SEP-2000; 2000US-0236327P.
PR 29-SEP-2000; 2000US-0236367P.
PR 29-SEP-2000; 2000US-0236368P.
PR 29-SEP-2000; 2000US-0236369P.
PR 29-SEP-2000; 2000US-0236370P.
PR 02-OCT-2000; 2000US-0236802P.
PR 02-OCT-2000; 2000US-0237037P.
PR 02-OCT-2000; 2000US-0237038P.
PR 02-OCT-2000; 2000US-0237039P.
PR 02-OCT-2000; 2000US-0237040P.
PR 13-OCT-2000; 2000US-0239935P.
PR 13-OCT-2000; 2000US-0239937P.
PR 20-OCT-2000; 2000US-0240960P.
PR 20-OCT-2000; 2000US-0241221P.
PR 20-OCT-2000; 2000US-0241785P.
PR 20-OCT-2000; 2000US-0241786P.
PR 20-OCT-2000; 2000US-0241787P.
PR 20-OCT-2000; 2000US-0241808P.
PR 20-OCT-2000; 2000US-0241809P.
PR 20-OCT-2000; 2000US-0241826P.
PR 01-NOV-2000; 2000US-0244617P.
PR 08-NOV-2000; 2000US-0246474P.
PR 08-NOV-2000; 2000US-0246475P.
PR 08-NOV-2000; 2000US-0246476P.
PR 08-NOV-2000; 2000US-0246477P.
PR 08-NOV-2000; 2000US-0246478P.
PR 08-NOV-2000; 2000US-0246523P.
PR 08-NOV-2000; 2000US-0246524P.
PR 08-NOV-2000; 2000US-0246525P.
PR 08-NOV-2000; 2000US-0246526P.
PR 08-NOV-2000; 2000US-0246527P.
PR 08-NOV-2000; 2000US-0246528P.
PR 08-NOV-2000; 2000US-0246532P.
PR 08-NOV-2000; 2000US-0246609P.
PR 08-NOV-2000; 2000US-0246610P.
PR 08-NOV-2000; 2000US-0246611P.
PR 08-NOV-2000; 2000US-0246613P.
PR 17-NOV-2000; 2000US-0249207P.
PR 17-NOV-2000; 2000US-0249208P.
PR 17-NOV-2000; 2000US-0249209P.
PR 17-NOV-2000; 2000US-0249210P.
PR 17-NOV-2000; 2000US-0249211P.
PR 17-NOV-2000; 2000US-0249212P.

PR 17-NOV-2000; 2000US-0249213P.
PR 17-NOV-2000; 2000US-0249214P.
PR 17-NOV-2000; 2000US-0249215P.
PR 17-NOV-2000; 2000US-0249216P.
PR 17-NOV-2000; 2000US-0249217P.
PR 17-NOV-2000; 2000US-0249218P.
PR 17-NOV-2000; 2000US-0249244P.
PR 17-NOV-2000; 2000US-0249245P.
PR 17-NOV-2000; 2000US-0249264P.
PR 17-NOV-2000; 2000US-0249265P.
PR 17-NOV-2000; 2000US-0249297P.
PR 17-NOV-2000; 2000US-0249299P.
PR 17-NOV-2000; 2000US-0249300P.
PR 01-DEC-2000; 2000US-0250160P.
PR 01-DEC-2000; 2000US-0250391P.
PR 05-DEC-2000; 2000US-0251030P.
PR 05-DEC-2000; 2000US-0251988P.
PR 05-DEC-2000; 2000US-0256719P.
PR 06-DEC-2000; 2000US-0251479P.
PR 08-DEC-2000; 2000US-0251856P.
PR 08-DEC-2000; 2000US-0251868P.
PR 08-DEC-2000; 2000US-0251869P.
PR 08-DEC-2000; 2000US-0251989P.
PR 08-DEC-2000; 2000US-0251990P.
PR 11-DEC-2000; 2000US-0254097P.
PR 05-JAN-2001; 2001US-0259678P.

XX (HUMA-) HUMAN GENOME SCI INC.
XX Rosen CA, Barash SC, Ruben SM;
XX WPI; 2001-483426/52.
XX Nucleic acids encoding human immune/hematopoietic antigen polypeptides,
XX useful for preventing, diagnosing and/or treating cancers and metastasis.
XX Disclosure; SEQ ID NO 22094; 3071pp + Sequence Listing; English.
XX AAK54951 to AAK64702 encode the human immune/haematopoietic antigen (I)
XX amino acid sequences given in AAM82170 to AAM91921. (I) have cytostatic
XX activity, and can be used in gene therapy and vaccine production. (I)
XX proteins and polynucleotides may be used in the prevention, diagnosis and
XX treatment of diseases associated with inappropriate (I) expression. For
XX example, they may be used to treat disorders associated with decreased
XX expression by rectifying mutations or deletions in a patient's genome
XX that affect the activity of (I) by expressing inactive proteins or to
XX supplement the patients own production of (I). Additionally, (I)
XX polynucleotides may be used to produce the secreted (I), by inserting the
XX nucleic acids into a host cell and culturing the cell to express the
XX protein. (I) proteins and polynucleotides may be used to prevent,
XX diagnose and treat immune/haematopoietic-related diseases, especially
XX cancers and cancer metastases of haematopoietic-derived cells. AAK64703
XX to AAK87694 represent human immune/haematopoietic antigen genomic
XX sequences from the present invention. AAK54942 to AAK54950 and AAM82169
XX represent sequences used in the exemplification of the present invention
XX
XX Sequence 33147 BP; 8370 A; 7682 C; 8382 G; 8713 T; 0 U; 0 Other;

Query Match 20.4%; Score 364.2; DB 4; Length 33147;
Best Local Similarity 96.6%; Pred. NO. 6.8e-88;
Matches 372; Conservative 0; Mismatches 13; Indels 0; Gaps 0;

QY 746 CTGATGCCAAATCCAAAAGGAGCGGAGAGAAAGAAAGAGGACCAATATCTCAT 805
Db 8894 CAGATGCCAAATCCAAAAGGAGCGGAGAGAAAGAAAGGACCAATATCTCAT 8953
QY 806 TTGATGATGAGGAAGATGAGCAGAACTCTGGGACGCTGTTTAAAAAGACACCTGGGCGAG 865
Db 8954 TTGATGATGAGGAAGATGAGCAGAACTCTGGGACACATGTTTAAAAAGACACCTGGGCGAG 9013
QY 866 GGGAGAGCTCAGAGGCAACTCCGACGCTCTCTGTAATATCATGTCGCCCTTTGAAA 925
Db 9014 GGGAGAGCTCAGAGGCAACTCCGACGCTCTCTGTAATATCATGTCGCCCTTTGAAA 9073

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QY 926 GCCCTTGGCCCTAATCAATGAAAGTACAGCAGCAACTCATGGAATTTGATTTCCC 985  
 Db 9074 GCCCTTGGCCCAAACTCCAATGGAAGTACAGCAGCAACTCTGGAATTTGATTTCCC 9133  
 QY 986 TGTCTTTGACCGGAGTTGGGTACCAAGCTTGTATGAAAAGCATCATGATGAAG 1045  
 Db 9134 TGTCTTTGACCGGAGTTGGGTACCAAGCTTGTATGAAAAGCATCATGATGAAG 9193  
 QY 1046 ATGTGGATGAAAACGAAGATGACCTGTATGAAAACCTCATCAGGAAGGACACAGGGGCC 1105  
 Db 9194 ATGTGGATGAAAACGAAGATGACCTGTATGAAAACCTCATCAGGAAGGACACAGGGGCC 9253  
 QY 1106 ACTCGGAGTCCCGAGAAGCACT 1130  
 Db 9254 ACTCAGAGTCGCCCGAGAAGTAAGT 9278

RESULT 9  
 ABS72233 standard; cDNA; 280 BP.  
 XX  
 AC ABS72233;  
 XX  
 DT 03-DEC-2002 (first entry)  
 XX  
 DE Human gene trapped sequence (GTS) #193.  
 XX  
 KW Human; gene trapped sequence; GTS; gene; ss; cancer; autoimmune disease;  
 KW lupus; scleroderma; Crohn's disease; multiple sclerosis; immune disorder;  
 KW inflammatory bowel disease; schizophrenia; psychosis; osteoarthritis;  
 KW rheumatoid arthritis; hypertension; skin disorder; acne; eczema; asthma;  
 KW cardiovascular disease; Parkinson's disease; atherosclerosis; Alzheimer's disease;  
 KW viral infection; parasitic infection; fungal infection;  
 KW bacterial infection; forensic analysis; cellular differentiation.  
 XX  
 OS Homo sapiens.  
 XX  
 PN US2002095031-A1.  
 XX  
 PD 18-JUL-2002.  
 XX  
 PF 03-MAY-2000; 2000US-00563817.  
 XX  
 PR 04-MAY-1999; 99US-0132343P.  
 XX  
 PA (NEHL/) NEHLS M C.  
 PA (ZAMB/) ZAMBROWICZ B.  
 PA (SAND/) SANDS A T.  
 XX  
 PI Nehls MC, Zambrowicz B, Sands AT;  
 XX  
 DR WPI; 2002-656030/70.  
 XX  
 PT New isolated or purified human gene trapped sequences, useful for gene  
 PT discovery, as markers for gene expression analysis, identifying and  
 PT mapping the coding regions of human genome, or determining the genetic  
 PT basis of human disease.  
 XX  
 PS Claim 1; SEQ ID NO 201; 36pp; English.  
 XX  
 CC The invention relates to isolated or purified polynucleotides that  
 CC correspond to human gene trapped sequences (GTSs). The human GTSs are  
 CC useful for gene discovery and as markers for gene expression analysis,  
 CC for identifying and mapping the coding regions of the mammalian,  
 CC particularly human, genome, for forensic analysis, and for determining  
 CC the genetic basis of human disease. The peptides and proteins encoded by  
 CC the polynucleotides are useful for generating antibodies, as reagents in  
 CC diagnostic assays and in identifying other cellular gene products  
 CC involved in the regulation of development and cellular differentiation of  
 CC various cell types. The peptides are also useful as reagents in assays  
 CC for screening of compounds used in treating disorders affecting

CC development and cell differentiation. The GTSs are also useful in  
 CC treating or ameliorating diseases associated with the expression of  
 CC mutant or normal variants of the GTSs, e.g. cancer, autoimmune diseases,  
 CC lupus, scleroderma, Crohn's disease, multiple sclerosis, inflammatory  
 CC bowel disease, immune disorders, schizophrenia, psychosis, inflammatory  
 CC disorders, diabetes, skin disorders such as acne or eczema,  
 CC osteoarthritis, rheumatoid arthritis, hypertension, atherosclerosis,  
 CC cardiovascular diseases, Alzheimer's disease, Parkinson's disease,  
 CC osteoporosis, asthma, infertility, and viral, parasitic, fungal or  
 CC bacterial infections. This sequence represents a human GTS of the  
 CC invention  
 XX  
 SQ Sequence 280 BP; 71 A; 66 C; 91 G; 51 T; 0 U; 1 Other;  
 Query Match 15.5%; Score 275.8; DB 6; Length 280;  
 Best Local Similarity 98.9%; Pred. No. 9.2e-65;  
 Matches 277; Conservative 0; Mismatches 3; Indels 0; Gaps 0;  
 QY 1349 GTCTCTGAGCAGCCTTTACCTTCTGCCTCAGTGCAGAGTCCATCACAATTAGTGAAC 1408  
 Db 1 GGCTATGAGCAGCCTGNTACCTTCTGCCTCAGTGCAGAGTCCATCACAATTAGTGAAC 60  
 QY 1409 TGGCCAGGCCACTGTGCGCCATGATGAACAGGAAGGATGAGTGGAGGAGAGACAGAT 1468  
 Db 61 TGGCCAGGCCACTGTGCGCCATGATGAACAGGAAGGATGAGTGGAGGAGAGACAGAT 120  
 QY 1469 CACTGCGAAAACCTGCTCGACGGTGAGATGGAGCACTCAGCGCGCTCCGGCAAGAGGTGG 1528  
 Db 121 CACTGCGAAAACCTGCTCGACGGTGAGATGGAGCACTCAGCGCGCTCCGGCAAGAGGTGG 180  
 QY 1529 ACACCTTTGAAAAGAGGTTGGCTGAAACAGGAGGAGCGGCGGATCAAGTCCAGGCGC 1588  
 Db 181 ACACCTTTGAAAAGAGGTTGGCTGAAACAGGAGGAGCGGCGGATCAAGTCCAGGCGC 240  
 QY 1589 TGGCCAGCTATCTTTCCTATTTTGTGAGGAGATTCTAAC 1628  
 Db 241 TGGCCAGCTATCTTTCCTATTTTGTGAGGAGATTCTAAC 280

RESULT 10  
 AAS64410  
 ID AAS64410 standard; cDNA; 454 BP.  
 XX  
 AC AAS64410;  
 XX  
 DT 13-FEB-2002 (first entry)  
 XX  
 DE DNA encoding novel human diagnostic protein #214.  
 XX  
 KW Human; chromosome mapping; gene mapping; gene therapy; forensic;  
 KW food supplement; medical imaging; diagnostic; genetic disorder; ss.  
 XX  
 OS Homo sapiens.  
 XX  
 PN WO200175067-A2.  
 XX  
 PD 11-OCT-2001.  
 XX  
 PF 30-MAR-2001; 2001WO-US008631.  
 XX  
 PR 31-MAR-2000; 2000US-00540217.  
 PR 23-AUG-2000; 2000US-00649167.  
 XX  
 PA (HYSE-) HYSEQ INC.  
 XX  
 PI Drmanac RT, Liu C, Tang YT;  
 XX  
 DR WPI; 2001-639362/73.  
 DR P-PSDB; ABG00223.  
 XX  
 PT New isolated polynucleotide and encoded polypeptides, useful in  
 PT diagnostics, forensics, gene mapping, identification of mutations  
 PT responsible for genetic disorders or other traits and to assess

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PT XX  biodiversity.
PS XX  Claim 1; SEQ ID NO 214; 103pp; English.
XX
CC  The invention relates to isolated polynucleotide (I) and polypeptide (II)
CC  sequences. (I) is useful as hybridisation probes, polymerase chain
CC  reaction (PCR) primers, oligomers, and for chromosome and gene mapping,
CC  and in recombinant production of (II). The polynucleotides are also used
CC  in diagnostic as expressed sequence tags for identifying expressed
CC  genes. (I) is useful in gene therapy techniques to restore normal
CC  activity of (II) or to treat disease states involving (II). (II) is
CC  useful for generating antibodies against it, detecting or quantitating a
CC  polypeptide in tissue, as molecular weight markers and as a food
CC  supplement. (II) and its binding partners are useful in medical imaging
CC  of sites expressing (II). (I) and (II) are useful for treating disorders
CC  involving aberrant protein expression or biological activity. The
CC  polypeptide and polynucleotide sequences have applications in
CC  diagnostics, forensics, gene mapping, identification of mutations
CC  responsible for genetic disorders or other traits to assess biodiversity
CC  and to produce other types of data and products dependent on DNA and
CC  amino acid sequences. AAS64197-AAS94564 represent novel human diagnostic
CC  coding sequences of the invention. Note: The sequence data for this
CC  patent did not appear in the printed specification, but was obtained in
CC  electronic format directly from WIPO at
CC  ftp.wipo.int/pub/published_pct_sequences
XX
SQ  Sequence 454 BP; 107 A; 110 C; 151 G; 86 T; 0 U; 0 Other;
    Query Match      13.9%; Score 247; DB 5; Length 454;
    Best Local Similarity 100.0%; Pred. No. 8.7e-57;
    Matches 247; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
QY  1  ATAGCGGATCACAGAACCAATGACAAAGACAAATTTCTGCTGAGCGACTGCTGGATGCA 60
DB  68  ATGAGCGGATCACAGAACCAATGACAAAGACAAATTTCTGCTGAGCGACTGCTGGATGCA 127
QY  61  GTGAACAGTCCAGATCCGCTTTGAGGGAGAAAGAGATTGCTCGGATTCGACAGC 120
DB  128  GTGAACAGTCCAGATCCGCTTTGAGGGAGAAAGAGATTGCTCGGATTCGACAGC 187
QY  121  AGGCTCACCTGTCTGTGTCGCCAGTTTGAAGCCGTCTCGAGCATGGCTTGAAGAGAGT 180
DB  188  AGGCTCACCTGTCTGTGTCGCCAGTTTGAAGCCGTCTCGAGCATGGCTTGAAGAGAGT 247
QY  181  CGAGGATTTGGACTACAGCGGAGCGATCAAGCAGCGAGCGGGCTTTGCCAGCAAAACC 240
DB  248  CGAGGATTTGGACTACAGCGGAGCGATCAAGCAGCGAGCGGGCTTTGCCAGCAAAACC 307
QY  241  GAAACAG 247
DB  308  GAAACAG 314
XX
RESULT 11
ACH75347/C
ID  ACH75347 standard; DNA; 542 BP.
XX
AC  ACH75347;
XX
DT  29-JUL-2004 (first entry)
XX
DE  Human genome derived single exon probe #8542.
XX
KW  Human; probe; ss; gene expression; single exon probe; microarray;
KW  alternative splicing event; genomic alteration.
XX
OS  Homo sapiens.
XX
PN  US2003194704-A1.
XX
PD  16-OCT-2003.
XX
PP  03-APR-2002; 2002US-00029386.
XX

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XX PR  biodiversity.
XX PR  Claim 1; SEQ ID NO 214; 103pp; English.
XX
PA  (PENN/) PENN S G.
PA  (RANK/) RANK D R.
PA  (HANZ/) HANZEL D K.
XX
PI  Penn SG, Rank DR, Hanzel DK;
XX  WPI; 2004-119264/12.
XX
PT  New human genome-derived single exon nucleic acid probes useful for human
PT  gene expression analysis, for identifying or characterizing alternative
PT  splicing events, for assessing genomic alterations or as tools for
PT  surveying tissues.
XX
PS  Claim 15; SEQ ID NO 8542; 80pp; English.
XX
CC  The invention relates to a nucleic acid probe for measuring human gene
CC  expression, comprising any of the 27,400 fully defined nucleotide
CC  sequences in the specification, or their complements or fragments, and
CC  encoding at least 8 amino acids of any of the 6888 amino acid sequences
CC  fully defined in the specification. The probe is a single exon probe that
CC  hybridises under high stringency conditions to a nucleic acid molecule
CC  expressed in human cells or tissues. Also included are a spatially-
CC  addressable set of single exon nucleic acid probes for measuring human
CC  gene expression (comprising a plurality of single exon nucleic acid
CC  probes cited above, where each of the plurality of probes is separately
CC  and addressably isolatable or amplifiable from the plurality), a single
CC  exon microarray for measuring human gene expression, a method of
CC  measuring human gene expression, a vector comprising the single exon
CC  probe cited above, an ORF-encoded peptide comprising at least 8
CC  contiguous amino acids of any of the above-mentioned amino acid
CC  sequences (optionally with conservative amino acid substitutions), an
CC  isolated antibody that binds specifically to a peptide cited above,
CC  methods of selling and/or licensing single exon probes or microarrays to
CC  a customer desiring to measure gene expression, a method of providing
CC  human gene expression data by subscription, and a computer-readable
CC  storage medium which contains a database having a plurality of records
CC  (each record including data on the expression of a single exon probe
CC  cited above. The probe, methods and apparatus are useful in gene
CC  expression analysis. The probes may be used as tools for surveying
CC  tissues to detect the presence of expressed messages that contain their
CC  specific exon, or in constructing genome-derived single exon microarrays.
CC  In addition, the probes are used in identifying and characterising
CC  alternative splicing events, in detecting and characterising gross
CC  alterations in the genomic locus that includes their exon, in assessing
CC  smaller genomic alterations, in priming the synthesis of nucleic acids,
CC  or in expressing the ORF-encoded peptide. The present sequence is a human
CC  single exon probe of the invention. Note: The sequence data for this
CC  patent did not form part of the printed specification, but was obtained
CC  in electronic format directly from USPTO at
CC  seqdata.uspto.gov/sequence.html?DocID=20030194704
XX
SQ  Sequence 542 BP; 135 A; 143 C; 129 G; 135 T; 0 U; 0 Other;
    Query Match      12.8%; Score 227.8; DB 12; Length 542;
    Best Local Similarity 96.1%; Pred. No. 1.7e-51;
    Matches 245; Conservative 0; Mismatches 7; Indels 3; Gaps 1;
XX
QY  497  CAGGCTGTAACCTCATCTCTTTGGCGATTAAACATGACACAAAGGATTTCGAACGGGCGAGA 556
DB  495  CAGGCTGTAACCTCATCTCTTTGGCGATTAAACATGACACAAAGGATTTCGAACGGGCGAGA 436
QY  557  GTAAGTTTGTCTCCACCGTTTCAGACCTCTTAAAGGAGTCAACGCGAAGACCTGCTCT 616
DB  435  GTAAGTTTGTCTCCACCGTTTCAGACCTCTTAAAGGAGTCAACGCGAAGTGTGA---CCT 379
QY  617  TGCTGAAGGAGTCCAGCAAGGAGTGAAGCCCTGTTTCAGGAGATCACAGCTCTCTCTG 676
DB  378  TGCTGAAGGAGTCCAGCAAGGAGTGAAGCCCTGTTTCAGGAGATCACAGCTCTCTCTG 319
QY  677  CCGTCTCCATCTCATCAAAACCTGAACAGGAGACCGACCCCTTGCCTGTGTCGTCCAGGA 736

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Db 318 CCATCTCATCTCATCAAACTGAACAGGAGACCGCCCTTCCCGTCTGTCCAGGA 259  
QY 737 ATGTCAGTCTGTGATG 751  
Db 258 ATGTCAGTCTGTGATG 244  
RESULT 12  
ACH89047/C  
ID ACH89047 standard; DNA; 284 BP.  
XX  
AC ACH89047;  
DT 29-JUL-2004 (first entry)  
XX Human genome derived single exon probe #22242.  
XX Human; probe; ss; gene expression; single exon probe; microarray;  
KW alternative splicing event; genomic alteration.  
XX Homo sapiens.  
XX US2003194704-A1.  
XX 16-OCT-2003.  
XX 03-APR-2002; 2002US-00029386.  
XX 03-APR-2002; 2002US-00029386.  
XX (PENN/) PENN S G.  
PA (RANK/) RANK D R.  
PA (HANZ/) HANZEL D K.  
XX Penn SG, Rank DR, Hanzel DK;  
XX WPI; 2004-119264/12.  
XX New human genome-derived single exon nucleic acid probes useful for human  
PT gene expression analysis, for identifying or characterizing alternative  
PT splicing events, for assessing genomic alterations or as tools for  
PT surveying tissues.  
XX Claim 1; SEQ ID NO 22242; 80pp; English.  
XX The invention relates to a nucleic acid probe for measuring human gene  
CC expression, comprising any of the 27,400 fully defined nucleotide  
CC sequences in the specification, or their complements or fragments, and  
CC encoding at least 8 amino acids of any of the 688 amino acid sequences  
CC fully defined in the specification. The probe is a single exon probe that  
CC hybridizes under high stringency conditions to a nucleic acid molecule  
CC expressed in human cells or tissues. Also included are a spatially-  
CC addressable set of single exon nucleic acid probes for measuring human  
CC gene expression (comprising a plurality of single exon nucleic acid  
CC probes cited above, where each of the plurality of probes is separately  
CC and addressably isolatable or amplifiable from the plurality), a single  
CC exon microarray for measuring human gene expression, a method of  
CC measuring human gene expression, a vector comprising the single exon  
CC probe cited above, an ORF-encoded peptide comprising at least 8  
CC contiguous amino acids of any of the above-mentioned amino acid  
CC sequences (optionally with conservative amino acid substitutions), an  
CC isolated antibody that binds specifically to a peptide cited above,  
CC a method of selling and/or licensing single exon probes or microarrays to  
CC a customer desiring to measure gene expression, a method of providing  
CC human gene expression data by subscription, and a computer-readable  
CC storage medium which contains a database having a plurality of records  
CC (each record including data on the expression of a single exon probe  
CC cited above). The probe, methods and apparatus are useful in gene  
CC expression analysis. The probes may be used as tools for surveying  
CC tissues to detect the presence of expressed messages that contain their  
CC specific exon, or in constructing genome-derived single exon microarrays.  
CC In addition, the probes are used in identifying and characterising

CC alternative splicing events, in detecting and characterising gross  
CC alterations in the genomic locus that includes their exon, in assessing  
CC smaller genomic alterations, in priming the synthesis of nucleic acids,  
CC or in expressing the ORF-encoded peptide. The present sequence is a human  
CC single exon probe of the invention. Note: The sequence data for this  
CC patent did not form part of the printed specification, but was obtained  
CC in electronic format directly from USPTO at  
CC seqdata.uspto.gov/sequence.html?DocID=20030194704  
XX  
SQ Sequence 284 BP; 60 A; 78 C; 74 G; 72 T; 0 U; 0 Other;  
Query Match 12.6%; Score 224.8; DB 12; Length 284;  
Best Local Similarity 96.0%; Pred. No. 8e-51;  
Matches 242; Conservative 0; Mismatches 7; Indels 3; Gaps 1;  
QY 500 GTCTGAATCCATACTCTTTGCGATTAAACATCGACACAAAGGATTTTCAACGGGCAGAGTA 559  
Db 284 GTCCGAATCTCATCTCTTTGCGATTAAACATCGACACAAAGGATTTTCAACGGGCAGAGTA 225  
QY 560 AGTTTGTCTCCACCGTTTTCAGACCTCTTAAAGGAGTCAACGCGAGAGTCACTCTTGC 619  
Db 224 AGTTTGTCTCCACCGTTTTCAGACCTCTTAAAGGAGTCAACGCGAGAGTCACTCTTGC 168  
QY 620 TGAAGGAGTCCAGCAAGGAGTGTGAGCAGCTGTTCAGGAGATCACAGCCTCTCTGCCG 679  
Db 167 TGAAGGAGTCCAGCAAGGAGTGTGAGCAGCTGTTCAGGAGATCACAGCCTCTCTGCCA 108  
QY 680 TCTCCATCTCTCAACACCTGAAACAGGAGACCGACCCCTTGCCTGTCTGTCAGGAATG 739  
Db 107 TCTCCATCTCTCAACACCTGAAACAGGAGACCGACCCCTTGCCTGTCTGTCAGGAATG 48  
QY 740 TCAGTGTCTGATG 751  
Db 47 TCAGTGTCTGATG 36  
RESULT 13  
AAS92200  
ID AAS92200 standard; cDNA; 463 BP.  
XX  
AC AAS92200;  
XX  
DT 13-FEB-2002 (first entry)  
XX  
XX DNA encoding novel human diagnostic protein #28004.  
XX Human; chromosome mapping; gene mapping; gene therapy; forensic;  
KW food supplement; medical imaging; diagnostic; genetic disorder; ss.  
XX Homo sapiens.  
XX WO200175067-A2.  
XX  
PD 11-OCT-2001.  
XX  
XX 30-MAR-2001; 2001WO-US008631.  
XX  
XX 31-MAR-2000; 2000US-00540217.  
PR 23-AUG-2000; 2000US-00649167.  
XX  
XX (HYSE-) HYSEQ INC.  
PA  
XX Drmanac RT, Liu C, Tang YT;  
XX  
XX WPI; 2001-639362/73.  
DR P-PSDB; ABG28013.  
XX  
XX New isolated polynucleotide and encoded polypeptides, useful in  
PT diagnostics, forensics, gene mapping, identification of mutations  
PT responsible for genetic disorders or other traits and to assess  
PT biodiversity.  
XX  
PS Claim 1; SEQ ID NO 28004; 103pp; English.

XX The invention relates to isolated polynucleotide (I) and polypeptide (II)  
 CC sequences. (I) is useful as hybridisation probes, polymerase chain  
 CC reaction (PCR) primers, oligomers, and for chromosome and gene mapping,  
 CC and in recombinant production of (II). The polynucleotides are also used  
 CC in diagnostics as expressed sequence tags for identifying expressed  
 CC genes. (I) is useful in gene therapy techniques to restore normal  
 CC genon. (I) is useful in gene therapy techniques to restore normal  
 CC activity of (II) or to treat disease states involving (II). (II) is  
 CC useful for generating antibodies against it, detecting or quantitating a  
 CC polypeptide in tissue, as molecular weight markers and as a food  
 CC supplement. (II) and its binding partners are useful in medical imaging  
 CC of sites expressing (II). (I) and (II) are useful for treating disorders  
 CC involving aberrant protein expression or biological actions in  
 CC polypeptide and polynucleotide sequences have applications in  
 CC diagnostics, forensics, gene mapping, identification of mutations  
 CC responsible for genetic disorders or other traits to assess biodiversity  
 CC and to produce other types of data and products dependent on DNA and  
 CC amino acid sequences. AAS64197-AAS94564 represent novel human diagnostic  
 CC coding sequences. AAS64197-AAS94564 represent novel human diagnostic  
 CC coding sequences of the invention. Note: The sequence data for this  
 CC patent did not appear in the printed specification, but was obtained in  
 CC electronic format directly from WIPO at  
 CC ftp.wipo.int/pub/published\_pct\_sequences  
 XX  
 SQ Sequence 463 BP; 124 A; 133 C; 122 G; 84 T; 0 U; 0 Other;  
 Query Match 11.6%; Score 207.6; DB 5; Length 463;  
 Best Local Similarity 93.9%; Pred. No. 5.1e-46;  
 Matches 216; Conservative 0; Mismatches 14; Indels 0; Gaps 0;  
 QY 1219 GGCGTGGCTCTACAGCCAGCAGATGCCCCCTCGGAGCTCGAGACGGGACAGGA 1278  
 Db GTCTCAGCCCTGGCTGAAACAAAGATGCCCCCTCGGAGCTCGAGACGGGACAGGA 293  
 QY 1279 CCAGAGGACCACTTCTCCCGGATCCTGGACTTCGGTACAGTGTGGAAGCCAGCTCTCCA 1338  
 Db CCAGAGGACCACTTCTCCCGGATCCTGGACTTCGGTACAGTGTGGAAGCCAGCTCTCCA 353  
 QY 1339 GGCACGGAGTCTCTGAGCAGCCTGTACCTTCTGCTTCAGTGCAGAGTCCATGACA 1398  
 Db GGCACGGAGTCTCTGAGCAGCCTGTACCTTCTGCTTCAGTGCAGAGTCCATGACA 413  
 QY 1399 ATTAGTGAAGTGGCCAGGCGCACTGTGGCCATGATGAACAGGAAGATGA 1448  
 Db ATTAGTGAAGTGGCCAGGCGCACTGTGGCCATGATGAACAGGAAGATGA 463  
 RESULT 14  
 AAS73549/c  
 ID AAS73549 standard; cDNA; 474 BP.  
 XX  
 AC AAS73549;  
 XX  
 XX  
 DT 13-FEB-2002 (first entry)  
 XX  
 DE DNA encoding novel human diagnostic protein #9353.  
 XX  
 XX Human; chromosome mapping; gene mapping; gene therapy; forensic;  
 KW food supplement; medical imaging; diagnostic; genetic disorder; ss.  
 XX Homo sapiens.  
 OS  
 XX  
 PN WO200175067-A2.  
 XX  
 PD 11-OCT-2001.  
 XX  
 PF 30-MAR-2001; 2001WO-US008631.  
 XX  
 PF 31-MAR-2000; 2000US-00540217.  
 PR 23-AUG-2000; 2000US-00649167.  
 XX  
 XX (HYSB-) HYSBQ INC.  
 PA  
 XX Drmanac RT, Liu C, Tang YT;

XX  
 DR WPI; 2001-639362/73.  
 DR P-PSDB; ABG09362.  
 XX  
 PT New isolated polynucleotide and encoded polypeptides, useful in  
 PT diagnostics, forensics, gene mapping, identification of mutations  
 PT responsible for genetic disorders or other traits and to assess  
 PT biodiversity.  
 XX  
 PS Claim 1; SEQ ID NO 9353; 103pp; English.  
 XX  
 CC The invention relates to isolated polynucleotide (I) and polypeptide (II)  
 CC sequences. (I) is useful as hybridisation probes, polymerase chain  
 CC reaction (PCR) primers, oligomers, and for chromosome and gene mapping,  
 CC and in recombinant production of (II). The polynucleotides are also used  
 CC in diagnostics as expressed sequence tags for identifying expressed  
 CC genes. (I) is useful in gene therapy techniques to restore normal  
 CC activity of (II) or to treat disease states involving (II). (II) is  
 CC useful for generating antibodies against it, detecting or quantitating a  
 CC polypeptide in tissue, as molecular weight markers and as a food  
 CC supplement. (II) and its binding partners are useful in medical imaging  
 CC of sites expressing (II). (I) and (II) are useful for treating disorders  
 CC involving aberrant protein expression or biological actions in  
 CC polypeptide and polynucleotide sequences have applications in  
 CC diagnostics, forensics, gene mapping, identification of mutations  
 CC responsible for genetic disorders or other traits to assess biodiversity  
 CC and to produce other types of data and products dependent on DNA and  
 CC amino acid sequences. AAS64197-AAS94564 represent novel human diagnostic  
 CC coding sequences of the invention. Note: The sequence data for this  
 CC patent did not appear in the printed specification, but was obtained in  
 CC electronic format directly from WIPO at  
 CC ftp.wipo.int/pub/published\_pct\_sequences  
 XX  
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 DT 06-OCT-2000 (first entry)  
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 KW gene therapy; chromosome mapping; ss.  
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OS Homo sapiens.
PN EP1033401-A2.
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PD 06-SEP-2000.
XX
XX 21-FEB-2000; 2000EP-00200610.
XX
XX 26-FEB-1999; 99US-0122487P.
XX
XX (GBST ) GENSET.
XX
XX Dumas Milne Edwards J, Duclert A, Giordano J;
XX WPI; 2000-500381/45.
XX
XX New nucleic acid that is a 5' expressed sequence tag (5' EST) for
PT obtaining cDNAs and genomic DNAs that correspond to 5'ESTs and for
PT diagnostic, forensic, gene therapy and chromosome mapping procedures.
XX
XX Claim 1; SEQ ID NO 16002; 71pp + Sequence Listing; English.
XX
XX The present sequence is one of a large number of 5' ESTs derived from
CC mRNAs encoding secreted proteins. No ORF has yet been conclusively
CC identified within the present sequence. The 5' ESTs were prepared from
CC total human RNAs or polyA+ RNAs derived from 30 different tissues. EST
CC sequences usually correspond mainly to the 3' untranslated region (UTR)
CC of the mRNA because they are often obtained from oligo-dT primed cDNA
CC libraries. Such ESTs are not well suited for isolating cDNA sequences
CC derived from the 5' ends of mRNAs and even in those cases where longer
CC cDNA sequences have been obtained, the full 5' UTR is rarely included. 5'
CC ESTs are derived from mRNAs with intact 5' ends and can therefore be used
CC to obtain full length cDNAs and genomic DNAs. 5' ESTs are also used in
CC diagnostic, forensic, gene therapy and chromosome mapping procedures.
CC They are used to obtain upstream regulatory sequences and to design
CC expression and secretion vectors
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Best Local Similarity 99.3%; Pred. No. 6.2e-26;
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DEFINITION Sequence 6 from Patent WO0116314.
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VERSION AX089606.1 GI:13443798
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Homo sapiens
REFERENCE 1
AUTHORS Wallach,D., Malinin,N., Sinha,I.W. and Leu,S.
TITLE Iren protein, its preparation and use
JOURNAL Patent: WO 0116314-A 6 08-MAR-2001;
YEDA RESEARCH AND DEVELOPMENT Co. LTD. (IL)
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VERSION AX089605.1 GI:13443797  
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ORGANISM Homo sapiens  
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REFERENCE 1  
AUTHORS Wallach,D., Malinin,N., Sinha,I.W. and Leu,S.  
TITLE Iren protein, its preparation and use  
JOURNAL Patent: WO 0116314-A 5 08-MAR-2001;  
YEDA RESEARCH AND DEVELOPMENT Co. LTD. (IL)  
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RESULT 5  
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LOCUS BD157657 2248 bp DNA linear PAT 17-JAN-2003  
DEFINITION Primer for synthesizing full-length cDNA and use thereof.  
ACCESSION BD157657  
VERSION BD157657.1 GI:27863415  
KEYWORDS JP 2002191363-A/12500.  
SOURCE Homo sapiens (human)  
ORGANISM Homo sapiens  
Bukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
1 (bases 1 to 2248)  
Ota,T., Isogai,T., Nishikawa,T., Hayashi,K., Saito,K., Yamamoto,J.,  
Ishii,S., Sugiyama,T., Wakamatsu,A., Nagai,K. and Otsuki,T.  
Primer for synthesizing full-length cDNA and use thereof  
Patent: JP 2002191363-A 12500 09-JUL-2002;  
HELIIX RESEARCH INSTITUTE  
OS Homo sapiens (human)  
PN JP 2002191363-A/12500  
PD 09-JUL-2002  
PF 28-JUL-2000 JP 2000280990  
PI TOSHIO OTA,TAKAO ISOGAI,TETSUO NISHIKAWA,KOJI HAYASHI,KAORU  
PI SAITO,  
PI JUNICHI YAMAMOTO,SHIZUKO ISHII,TOMOYASU SUGIYAMA,AI WAKAMATSU,  
PI KEIICHI NAGAI,TETSUJI OTSUKI  
PC C12N15/09,C07K14/47,C07K16/18,C12N1/15,C12N1/19,C12N1/21,C12N5/PC  
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FEATURES  
Query Match 63.4%; Score 1129.8; DB 6; Length 2248;  
Best Local Similarity 95.3%; Pred. No. 3.9e-267;  
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121 AGGTCACCTGTCTGTGTGCCCGCTTGAAGCCGCTCTCGAGCATGCTTGAAGAGGAGT 180  
183 AGGTCACCTGTCTGTGTGCCCGCTTGAAGCCGCTCTCGAGCATGCTTGAAGAGGAGT 242  
181 CGAGGATGGCACTCACAGCGGACGATCAAGCAGCAGCGGCTTTGCCAGCAAAACC 240

Db	243	CGAGGATTGGCACTCAGACGGCAGCGATCAAGCAGGACGGGCTTTGCCAGCAAAACC	302
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Db	303	GAACAGAGCCCTGTTCTGGTACTACGTGAAGAGGTCTCTCAACAGCACGAGCTGCAG	362
Qy	301	CGCTTCTACTCCCTGGCCACATCGCTCAGAGCTGGCGGGGCTCGCGCTGGCTGGCC	360
Db	363	CGCTTCTACTCCCTGGCCACATCGCTCAGAGCTGGCGGGGCTCGCGCTGGCTGGCC	422
Qy	361	TGTCCCTCAACGAACACTCCCTGGAGCGCTACTCTGCACATGCTCTCTGGCCGACCGCTGC	420
Db	423	TGTCCCTCAACGAACACTCCCTGGAGCGCTACTCTGCACATGCTCTCTGGCCGACCGCTGC	482
Qy	421	AGCTGAGCACTTTTATGAAGACTGGTCTTTTGTGATGATGAAGAGGTCCAGTATG	480
Db	483	AGCTGAGCACTTTTATGAAGACTGGTCTTTTGTGATGATGAAGAGGTCCAGTATG	542
Qy	481	CTTCTACCATGCGCAGAGTCTGAATCCATCTCTTTGCGATTAAACATCGACAACAG	540
Db	543	CTTCTACCATGCGCAGAGTCTGAATCCATCTCTTTGCGATTAAACATCGACAACAG	602
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Db	603	GATTGGAACGGGAGAGTAAGTTTGTCTCCACCGTTTTCAGACCTCTTAAAGAGTCAACG	662
Qy	601	CAGAACGTGACCTCTTGTGAAGAGTCCACGCAAGAGTGAAGCAGCTGTTTCAAGGAG	660
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Qy	661	ATCAGAGCTCTCTCGCTCTCCATCTCATCAACCTTCAACAGGAGACCGACCTCTTG	720
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Qy	721	CTGTCTGTCCAGGAATGTCAGTGTGTGATGCAAAATGCAAAAGAGCGGAGAAAGAA	780
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Qy	841	GTGTTTAAAGACACCTGGGCGAGGAGGCTCAGAGGACAACTCCGACCGCTCTCT	900
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Qy	901	GTCAATATCATGTCCGCTTTGAAAGCCCTTCGGGCTTAACCTCAATGGAAGTCAGAGC	960
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Qy	1141	ACCTGCTCTCCAGATGACAGCTGGGCTCCGTGAAGGTGTCGCAATGATCTCGAC	1200
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DEFINITION	Sequence 14016 from Patent EP1074617.				
ACCESSION	AX879111				
VERSION	AX879111.1				
KEYWORDS	GI:40033847				
SOURCE	Homo sapiens (human)				
ORGANISM	Homo sapiens				
REFERENCE	Ota, T., Isegai, T., Nishikawa, T., Hayashi, K., Saito, K., Yamamoto, J., Ishii, S., Sugiyama, T., Wakamatsu, A., Nagai, K. and Otsuki, T. Primers for synthesizing full-length cDNA and their use Patent: EP 1074617-A 14016 07-FEB-2001; (JP) Research Association for Biotechnology (JP) Location/Qualifiers				
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ORIGIN	Query Match 63.4%; Score 1129.8; DB 6; Length 2248;				
	Best Local Similarity 95.3%; Pred. No. 3.8e-267;				
	Matches 1164; Conservative 0; Mismatches 57; Indels 0; Gaps 0;				
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Qy	61	GTGAAACAGTGCAGATCGCTTTGGAGGAGAAAGAGATTGCCTCGATTCCGACAGC	120		
Db	123	GTGAAACAGTGCAGATCGCTTTGGAGGAGAAAGAGATTGCCTCGATTCCGACAGC	182		
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Db	183	AGGTCACCTGTCTGTGTCGCCAGTTTGAAGCCGCTCTGCAGCATGGCTTGAAGAGAGT	242		
Qy	181	CGAGGATTGGCACTCAGCGGGCAGCGATCAGCAGGAGCGGGCTTTGCCAGCAAAACC	240		
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Qy	241	GAAACAGAGCCCGTGTCTGTTACTACGTGAAGAGGTCTCTCAACAGCACGAGCTGCAG	300		
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Qy	301	CGCTTCTACTCCCTGCGCCACATCGCTCAGACGTGGCGGGGCTCGCGCTGGCTGGCG	360		
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Qy	361	TGTGCCCTCAACGAACACTCCCTGGAGCGCTACTGCAATGCTCTCTGGCCGACCGTGC	420		
Db	423	TGTGCCCTCAACGAACACTCCCTGGAGCGCTACTGCAATGCTCTCTGGCCGACCGTGC	482		
Qy	421	AGGCTGAGCACTTTTATGAAGACTGGTCTTTTGTGATGATGAAGAGGTCCAGTATG	480		
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Qy	481	CTTCTACCATGCGCAGAGTCTGAATCCATCTCTTTGCGATTAAACATCGACAACAG	540		
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 DEFINITION  
 ACCESSION AK022425  
 VERSION AK022425.1 GI:10433818  
 KEYWORDS oligo capping; fis (full insert sequence).  
 SOURCE Homo sapiens (human)  
 ORGANISM Homo sapiens  
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 REFERENCE 1  
 Ota, T., Suzuki, Y., Nishikawa, T., Otsuki, T., Sugiyama, T., Irie, R., Wakamatsu, A., Hayaishi, K., Sato, H., Nagai, K., Kimura, K., Makita, H., Sekine, M., Ohayashi, M., Nishi, T., Shibahara, T., Tanaka, T., Ishii, S., Yamamoto, J., Saito, K., Kawai, Y., Isono, Y., Nakamura, Y., Nagahara, K., Murakami, K., Yaeuda, T., Iwayanagi, T., Wagatsuma, M., Shiratori, A., Sudo, H., Hosoiri, T., Kaku, Y., Kodaira, H., Kondo, H., Sugawara, M., Takahashi, M., Kanda, K., Yokoi, T., Furuya, T., Kikkawa, E., Omura, Y., Abe, K., Kamihara, K., Katsuta, N., Sato, K., Tanikawa, M., Yanazaki, M., Ninomiya, K., Ishibashi, T., Yamashita, H., Murakawa, K., Fujimori, K., Tanai, H., Kimata, M., Watanabe, M., Hiraoka, S., Chiba, Y., Ishida, S., Ono, Y., Takiguchi, S., Watanabe, S.,

Yosida, M., Hotuta, T., Kusano, J., Kanehori, K., Takahashi-Fujii, A., Hara, H., Tanase, T., Nomura, Y., Togiya, S., Komai, F., Hara, R., Takeuchi, K., Arita, M., Imase, N., Mueashino, K., Yuuki, H., Oshima, A., Sasaki, N., Aotsuka, S., Yoshikawa, Y., Matsunawa, H., Ichihara, T., Shionata, N., Sano, S., Moriya, S., Momiyama, H., Satoh, N., Takami, S., Terahima, Y., Suzuki, O., Nakagawa, S., Senoh, A., Mizoguchi, H., Goto, Y., Shimizu, F., Wakebe, H., Hishigaki, H., Watanabe, T., Sugiyama, A., Takemoto, M., Kawakami, B., Yamazaki, M., Watanabe, K., Kumagai, A., Itakura, S., Fukuzumi, Y., Fujimori, Y., Komiyama, M., Tashiro, H., Tanigami, A., Fujiwara, T., Ono, T., Yanada, K., Fujii, Y., Ozaki, K., Hirao, M., Ohmori, Y., Kawabata, A., Hikiji, T., Kobatake, N., Inagaki, S., Ikema, Y., Okamoto, S., Okitani, R., Kawakami, T., Noguchi, H., Itoh, Y., Shigetani, K., Senba, T., Matsumura, K., Nakajima, Y., Mizuno, T., Morinaga, M., Sasaki, M., Togashi, T., Oyama, M., Hata, H., Watanabe, M., Komatsu, T., Mizushima-Sugano, J., Satoh, T., Shirai, Y., Takahashi, Y., Nakagawa, K., Okumura, K., Nagase, T., Nomura, N., Kikuchi, H., Masuho, Y., Yamashita, R., Nakamura, Y., Nigahara, K., Masuho, Y., Ninomiya, K., and Iwayanagi, T. NEDO human cDNA sequencing project

Unpublished  
 3 (bases 1 to 2248)  
 Isogai, T. and Otsuki, T.  
 Direct Submission  
 Submitted (23-AUG-2000) Takao Isogai, Helix Research Institute, Genomics Laboratory; 1532-3 Yana, Kisarazu, Chiba 292-0812, Japan (E-mail: genomics@hri.co.jp, Tel: 81-438-52-3975, Fax: 81-438-52-3986)  
 NEDO human cDNA sequencing project supported by Ministry of International Trade and Industry of Japan; cDNA full insert sequencing; Research Association for Biotechnology; cDNA library construction; 5'- & 3'-end one pass sequencing and clone selection; Helix Research Institute (supported by Japan Key Technology Center etc.) and Department of Virology, Institute of Medical Science, University of Tokyo.

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Query Match 63.4%; Score 1129.8; DB 9; Length 2248;  
 Best Local Similarity 95.3%; Pred. No. 3.8e-267;  
 Matches 1164; Conservative 0; Mismatches 57; Indels 0; Gaps 0;

ORIGIN  
 Query Match 63.4%; Score 1129.8; DB 9; Length 2248;  
 Best Local Similarity 95.3%; Pred. No. 3.8e-267;  
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RESULT 8
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DEFINITION Sequence 236 from Patent WO02068579.
ACCESSION CQ714292
VERSION CQ714292.1 GI:42275149
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SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1
AUTHORS Venter, C.J., Adams, M.C., Li, P.W. and Myers, E.W.
TITLE Kits, such as nucleic acid arrays, comprising a majority of
humanexons or transcripts, for detecting expression and other uses
thereof
JOURNAL Patent: WO 02068579-A 226 06-SEP-2002;
PE Corporation (NY) (US)
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LOCUS 6045 bp DNA linear PAT 17-JAN-2003
DEFINITION Primer for synthesizing full-length cDNA and use thereof.
ACCESSION BD160227
VERSION BD160227.1 GI:27865985
KEYWORDS JP 2002191363-A/15070.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE Mammalia; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Ota.T., Isogai.T., Nishikawa.T., Hayashi.K., Saito.K., Yamamoto.J.,
Ishii.S., Sugiyama.T., Wakamatsu.A., Nagai.K. and Otsuki.T.
TITLE Primer for synthesizing full-length cDNA and use thereof
JOURNAL Patent: JP 2002191363-A 15070 09-JUL-2002;
COMMENT HELIX RESEARCH INSTITUTE
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PN JP 2002191363-A/15070
PD 09-JUL-2002
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PI SAITO,
PI JUNICHI YAMAMOTO, SHIZUKO ISHII, TOMOYASU SUGIYAMA, AI WAKAMATSU,
PI KEIICHI NAGAI, TETSUJI OTSUKI
PC C12N15/09, C07K14/47, C07K16/18, C12N1/15, C12N1/19, C12N1/21, C12N5/10,
C12N5/11, C12N1/68, C12P21/08, G06F17/30, C12N15/00, C12N5/00 CC
PC, C12P21/02, C12Q1/68, C12P21/08, G06F17/30, C12N15/00, C12N5/00 CC
Primer for synthesizing full-length cDNA and use thereof FH Key
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FT CDS (364)..(954).
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Best Local Similarity 97.0%; Pred. No. 5.2e-190;
Matches 854; Conservative 0; Mismatches 22; Indels 4; Gaps 2;

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QY 305 TCTACTCCCTGCGCCACATCGCTCAGACGTGGCGCGGGTCCGGCTCGCTCGTGTG 364
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QY 365 CCCTCAACGAACACTCCCTGGAGCGGTACTCGACATGCTCTCGCGCGGACCGCTGAGGC 424
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QY 545 TGAACGGGCGAGAGTAAAGTTTGTCCACCGTTTCAGACCTCTTAAAGGAGTCAACCGAGA 604
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Ds	1052	CCCTCAACGAACTCTCCCTGGAGCGCTACCTGCACATGCTCTCTGGCCACCGCTGCAGGC	1111
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Ds	1769	CAGGAAGAGACACAGCGGCGCACTCGGAGTCCCGCGAGAA	1808
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LOCUS			
DEFINITION Homo sapiens cDNA FLJ13765 fis, clone PLACE400128, weakly similar			
to Mus musculus putative transcription factor mRNA.			
AK023827			
VERSION			
KEYWORDS			
SOURCE			
ORGANISM			
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;			
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.			

REFERENCE  
AUTHORS

1 Ota, T., Suzuki, Y., Nishikawa, T., Otsuki, T., Sugiyama, T., Irie, R., Wakamatsu, A., Hayashi, K., Sato, H., Nagai, K., Kimura, K., Makita, H., Sekine, M., Oobayashi, M., Nishi, T., Shibahara, T., Tanaka, K., Tanabe, T., Ishii, S., Yamamoto, J., Saito, K., Kawai, Y., Isono, Y., Nakamura, Y., Nagahara, K., Murakami, K., Yasuda, T., Iwayanagi, T., Wagatsuma, M., Shiratori, A., Sudo, H., Hosoiri, T., Kaku, Y., Kodaira, H., Kondo, H., Kikkawa, E., Omura, Y., Abe, K., Kamihara, K., Katsura, N., Sato, K., Tanikawa, M., Yamazaki, M., Ninomiya, K., Ishibashi, T., Yamashita, H., Murakawa, K., Fujimori, K., Tanai, H., Kimata, M., Watanabe, M., Hirakawa, S., Chiba, Y., Ishida, S., Ono, Y., Takiguchi, S., Watanabe, S., Yosida, M., Hotuta, T., Kusano, J., Kanehori, K., Takahashi, Fujii, A., Hara, H., Tanase, T., Nomura, Y., Togiya, S., Komai, F., Hara, R., Takeuchi, K., Arita, M., Imose, N., Musashino, K., Yuuki, H., Oshima, A., Sasaki, N., Aotsuka, S., Yoshihara, Y., Matsunawa, H., Tchihara, T., Shihata, N., Sano, S., Moriya, S., Momiyama, H., Satoh, N., Takami, S., Terashima, Y., Suzuki, O., Nakagawa, S., Senoh, A., Mizoguchi, H., Goto, Y., Shimizu, F., Wakebe, H., Hishigaki, H., Watanabe, T., Sugiyama, A., Takemoto, M., Kawakami, B., Yamazaki, M., Watanabe, K., Kumagai, A., Itakura, S., Fukuzumi, Y., Fujimori, Y., Komiyama, M., Tashiro, H., Tanigami, A., Fujiwara, T., Ono, T., Yanada, K., Fujii, Y., Ozaki, K., Hirao, M., Ohmori, Y., Kawabata, A., Hikiji, T., Kobatake, N., Inaguchi, H., Ikeda, Y., Okamoto, S., Okitani, R., Kawakami, T., Noguchi, S., Itoh, T., Shigeta, K., Senba, T., Matsunaga, K., Nakajima, Y., Mizuno, T., Morinaga, M., Sasaki, M., Togashi, T., Oyama, M., Hata, H., Watanabe, M., Komatsu, T., Mizushima-Sugano, J., Satoh, T., Shirai, Y., Takahashi, Y., Nakagawa, K., Okumura, K., Nagase, T., Nomura, N., Kikuchi, H., Masuho, Y., Yamashita, R., Nakai, K., Yada, T., Nakamura, Y., Ohara, O., Isogai, T. and Sugano, S. Complete sequencing and characterization of 21,243 full-length human cDNAs Nat. Genet. 36 (1), 40-45 (2004)

TITLE  
JOURNAL  
PUBMED  
REFERENCE  
AUTHORS

2 Isogai, T., Ota, T., Hayashi, K., Sugiyama, T., Otsuki, T., Suzuki, Y., Nishikawa, T., Nagai, K., Sugano, S., Takahashi-Fujii, A., Hara, H., Tanase, T., Nomura, Y., Togiya, S., Komai, F., Hara, R., Takeuchi, K., Arita, M., Nabekura, T., Ishii, S., Kawai, Y., Saito, K., Yamamoto, J., Wakamatsu, A., Nakamura, Y., Nagahara, K., Masuho, Y. and Oshima, A. NEDO human cDNA sequencing project Unpublished 3 (bases 1 to 6045) Isogai, T. and Otsuki, T. Direct Submission Submitted (21-AUG-2000) Takao Isogai, Helix Research Institute, Genomics Laboratory; 1532-3 Yana, Kisarazu, Chiba 292-0812, Japan (E-mail:genomics@hri.co.jp, Tel:81-438-52-3975, Fax:81-438-52-3986) NEDO human cDNA sequencing project supported by Ministry of International Trade and Industry of Japan; cDNA full insert sequencing: Research Association for Biotechnology; cDNA library construction, 5'- & 3'-end one pass sequencing and clone selection: Helix Research Institute (supported by Japan Key Technology Center etc.) and Department of Virology, Institute of Medical Science, University of Tokyo.

TITLE  
JOURNAL  
PUBMED  
REFERENCE  
AUTHORS  
JOURNAL

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FEATURES  
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DEFINITION
ACCESSION
VERSION
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Primer for synthesizing full-length cDNA and use thereof.
BD146965
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ACCESSION AX866903
VERSION AX866903.1 GI:40021252
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE 1
AUTHORS Ota,T., Isogai,T., Nishikawa,T., Hayaishi,K., Saito,K., Yamamoto,J.,
Ishii,S., Sugiyama,T., Wakamatsu,A., Nagai,K. and Otsuki,T.
TITLE Primers for synthesizing full-length cDNA and their use
JOURNAL Patent: EP 1074617-A 1808 07-FEB-2001;
Research Association for Biotechnology (JP)
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Best Local Similarity 99.0%; Pred. No. 2e-146;
Matches 663; Conservative 0; Mismatches 5; Indels 2; Gaps 2;
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